

=> d que

L1 16 SEA FILE=REGISTRY ABB=ON PLU=ON RRRPRPPYLPRRPP/SQSP
 L2 2 SEA FILE=REGISTRY ABB=ON PLU=ON ("PR 39"/CN OR "PR 39 (ION
 EXCHANGER)"/CN OR "PR 39 (PEPTIDE)"/CN)
 L4 53 SEA FILE=HCAPLUS ABB=ON PLU=ON L1 AND (L2 OR PR39 OR PR 39)
 L5 8645 SEA FILE=HCAPLUS ABB=ON PLU=ON ANGIOGENESIS+NT/CT
 L6 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L1 AND (L5 OR ANGIOGEN?)
 L7 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L4 AND (L5 OR ANGIOGEN?)
 L8 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L6 OR L7

=> d ibib abs hitstr l8 1-6

L8 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:220416 HCAPLUS

DOCUMENT NUMBER: 136:257252

TITLE: Method of modulating neovascularization

INVENTOR(S): Kovesdi, Imre

PATENT ASSIGNEE(S): Genvec, Inc., USA

SOURCE: PCT Int. Appl., 33 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002022176	A1	20020321	WO 2001-US28954	20010914
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2001091019	A5	20020326	AU 2001-91019	20010914
PRIORITY APPLN. INFO.:			US 2000-233001P	P 20000915
			WO 2001-US28954	W 20010914
AB	The present invention provides a method of modulating neovascularization in an animal. The method comprises administering to the animal two or more nucleic acid sequences, each nucleic acid sequence encoding at least one angiogenesis -modulation factor that acts upon a different angiogenic process, such that the nucleic acid sequences are expressed to produce the angiogenesis -modulation factors to modulate neovascularization in the animal. Modulating neovascularization includes the induction of neovascularization or, in the alternative, the inhibition or redn. of neovascularization.			
IT	139637-11-9, PR39 peptide			
	RL: BUU (Biological use, unclassified); PAC (Pharmacological activity); BIOL (Biological study); USES (Uses)			
	(method of modulating neovascularization)			
RN	139637-11-9 HCAPLUS			
CN	L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-			

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prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:319740 HCAPLUS

DOCUMENT NUMBER: 134:336214

TITLE: Method for **PR-39** peptide regulated stimulation of **angiogenesis**

INVENTOR(S): Simons, Michael; Gao, Youhe

PATENT ASSIGNEE(S): Beth Israel Deaconess Medical Center, USA

SOURCE: PCT Int. Appl., 52 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001030368	A1	20010503	WO 2000-US27552	20001006
W: AU, CA, JP				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

PRIORITY APPLN. INFO.: US 1999-426011 A 19991025

AB The present invention provides both a method and means for regulating **angiogenesis** within living cells, tissues, and organs in-situ. The regulation is performed using native **PR-39** peptide or one of its shorter-length homolog, for interaction with such proteasomes as one present in the cytoplasm of viable cells. The result of **PR-39** peptide interaction with proteasomes is a decrease in the intracellular degrdn. of active peptides such as HIF-1.alpha. and a consequential stimulation of **angiogenesis** in-situ.

IT 298702-64-4

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(**PR-39** peptide regulated stimulation of **angiogenesis**)

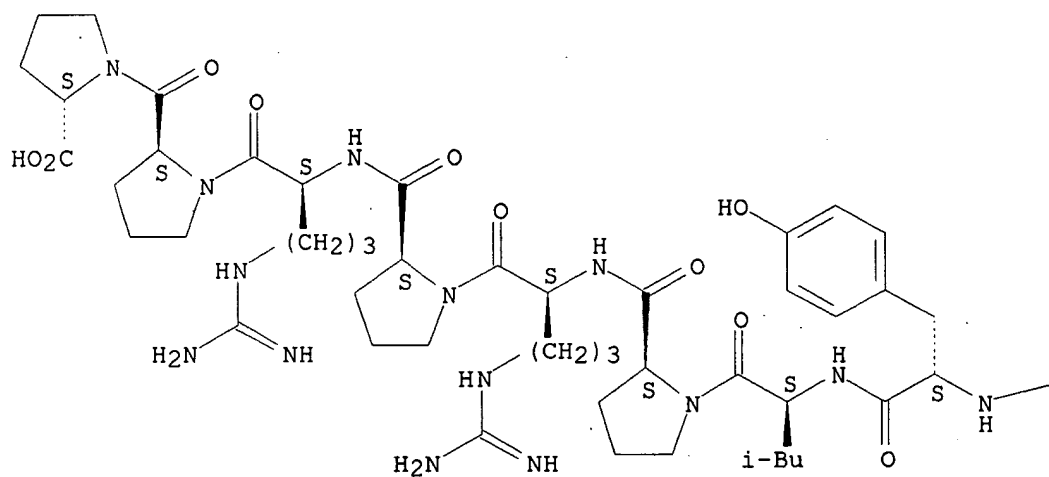
RN 298702-64-4 HCAPLUS

CN L-Proline, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl- (9CI) (CA INDEX NAME)

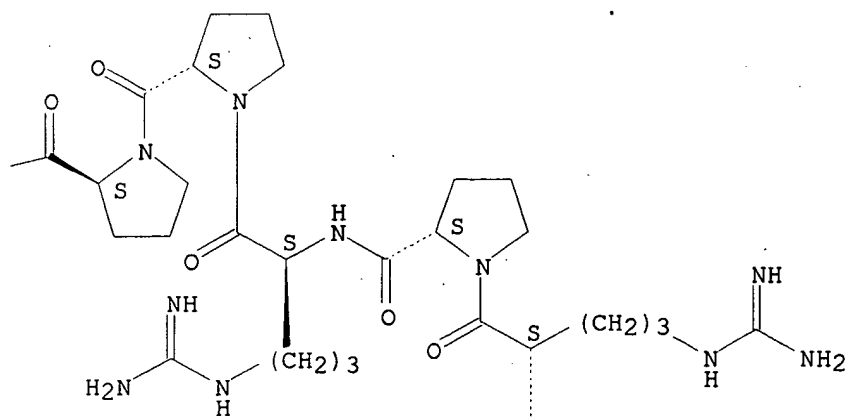
Absolute stereochemistry.

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PAGE 1-A



PAGE 1-B



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N=C(N)NCCSC(=O)NC(S)CNC(=O)NCCCNC(=N)N

Page 4

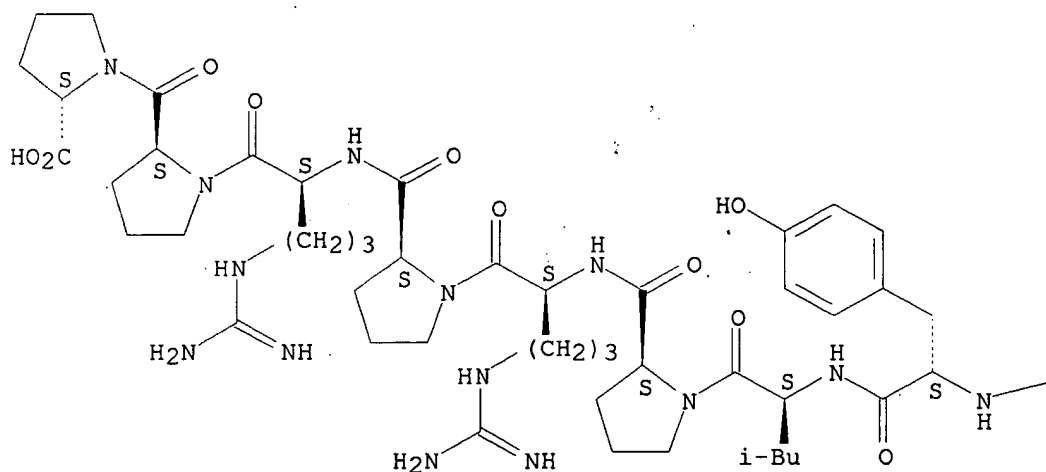
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IT 298702-64-4P

(PR-39 peptide-regulated stimulation of angiogenesis)

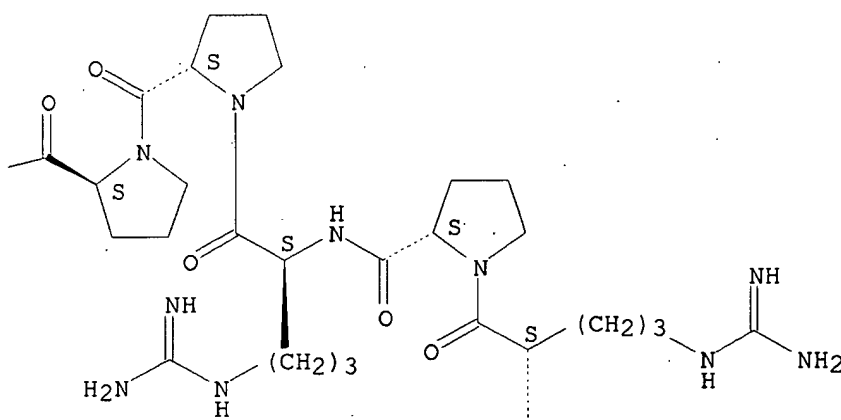
CN L-Proline, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-
(9CI) (CA INDEX NAME)

PAGE 1-A

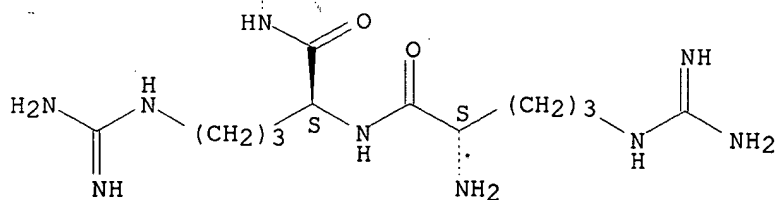


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PAGE 1-B



PAGE 2-B



IT 148046-54-2

RL: PRP (Properties)

(unclaimed protein sequence; method for PR-39
peptide regulated stimulation of angiogenesis)

RN 148046-54-2 HCAPLUS

CN L-Proline, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2000:178321 HCAPLUS

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DOCUMENT NUMBER: 133:205925
TITLE: **PR39**, a peptide regulator of **angiogenesis**. [Erratum to document cited in CA132:149677]
AUTHOR(S): Li, Jian; Post, Mark; Volk, Rudiger; Gao, Youhe; Li, Min; Metals, Caroline; Sato, Kaori; Tsai, Jo; Aird, William; Rosenberg, Robert D.; Hampton, Thomas G.; Li, Jianyi; Sellke, Frank; Carmeliet, Peter; Simons, Michael
CORPORATE SOURCE: Angiogenesis Research Center, Department of Surgery, Beth Israel Deaconess Medical Center and Harvard Medical School, Boston, MA, 02215, USA
SOURCE: Nature Medicine (New York) (2000), 6(3), 356
CODEN: NAMEFI; ISSN: 1078-8956
PUBLISHER: Nature America
DOCUMENT TYPE: Journal
LANGUAGE: English
AB The correct versions are given for Figs. 2a, c, and d on page 51; Fig. 3c on page 52; and Fig. 5b on page 53.
IT **139637-11-9, PR-39**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(**PR39** peptide in regulation of **angiogenesis** by inhibiting ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein (Erratum))
RN 139637-11-9 HCAPLUS
CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

L8 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 2000:46162 HCAPLUS
DOCUMENT NUMBER: 132:149677
TITLE: **PR39**, a peptide regulator of **angiogenesis**
AUTHOR(S): Li, Jian; Post, Mark; Volk, Rudiger; Gao, Youhe; Li, Min; Metals, Caroline; Sato, Kaori; Tsai, Jo; Aird, William; Rosenberg, Robert D.; Hampton, Thomas G.; Li, Jianyi; Sellke, Frank; Carmeliet, Peter; Simons, Michael
CORPORATE SOURCE: Angiogenesis Research Center, Department of Surgery both at Beth Israel Deaconess Medical Center and Harvard Medical School, Boston, MA, 02215, USA
SOURCE: Nature Medicine (New York) (2000), 6(1), 49-55
CODEN: NAMEFI; ISSN: 1078-8956
PUBLISHER: Nature America
DOCUMENT TYPE: Journal
LANGUAGE: English
AB Although tissue injury and inflammation are considered essential for the induction of **angiogenesis**, the mol. controls of this cascade are mostly unknown. Here we show that a macrophage-derived peptide,

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PR39, inhibited the ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein, resulting in accelerated formation of vascular structures in vitro and increased myocardial vasculature in mice. For the latter, coronary flow studies demonstrated that **PR39**-induced **angiogenesis** resulted in the prodn. of functional blood vessels. These findings show that **PR39** and related compds. can be used as potent inducers of **angiogenesis**, and that selective inhibition of hypoxia-inducible factor-1.alpha. degrdn. may underlie the mechanism of inflammation-induced **angiogenesis**.

IT 139637-11-9, **PR-39**

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(**PR39** peptide in regulation of **angiogenesis** by inhibiting ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein)

RN 139637-11-9 HCAPLUS

CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

REFERENCE COUNT: 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1996:357168 HCAPLUS

DOCUMENT NUMBER: 125:26311

TITLE: Synducin (syndecan expression-inducers) mediate modulation of tissue repair

INVENTOR(S): Gallo, Richard L.; Bernfield, Merton

PATENT ASSIGNEE(S): Children's Medical Center Corporation, USA

SOURCE: PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9609322	A2	19960328	WO 1995-US12080	19950922
WO 9609322	A3	19960523		
W: AU, CA, JP, KR				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5654273	A	19970805	US 1994-310722	19940922
AU 9538228	A1	19960409	AU 1995-38228	19950922
US 5863897	A	19990126	US 1996-728333	19961010
PRIORITY APPLN. INFO.:			US 1994-310722	19940922
			WO 1995-US12080	19950922

AB The membrane-permeating antibacterial peptide, **PR-39**, previously found only in the intestine, was purified from wound fluid and shown to possess syndecan-1 and syndecan-4 inductive activity specifically in mesenchymal cells. This is a newly recognized function that defines

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peptide contg. syndecan-inducing activity, and that are known as synducins. Therefore, a mol. with both antimicrobial and synducin activities is deposited in wounds where it can simultaneously reduce infection and the influence the action of growth factors, matrix components, and other cellular effectors involved in wound repair. Synducins, including **PR-39**, and derivs. thereof, as well as other proline and arginine-rich antimicrobial peptides, collectively referred to herein as "synducins", are therefore useful in the modulation of wound healing, as well as other disorders involving mesenchymal cells and cell surface mol. interaction, including metastatic disease, **angiogenesis**, restenosis, stasis or decubitus ulcers, and prevention of keloids.

IT 139637-11-9

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(synducins are syndecan expression-inducing peptides that mediate modulation of mesenchymal tissue repair)

RN 139637-11-9 HCAPLUS

CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolyl-glycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

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GenCore version 5.1.4 p5 4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 13, 2003, 10:32:27 ; Search time 35 Seconds
(without alignments)
57.107 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPRPPYLPRPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	90	100.0	15	21	PR-39 derived angi
2	90	100.0	15	22	Amino acid sequenc
3	90	100.0	15	22	PR-39 derived pept
4	90	100.0	19	17	Leukocyte O2- prod
5	90	100.0	26	17	Leukocyte O2- prod
6	90	100.0	26	19	Leukocyte O2- prod
7	90	100.0	39	14	proline/Arginine r
8	90	100.0	39	17	Antibacterial pept
9	90	100.0	39	17	Leukocyte O2- prod
10	90	100.0	39	17	Synducin peptide (
					Magainin-derived a

11	90	100.0	39	19	AAW75722	Proline/Arginine r
12	90	100.0	39	21	AAW75722	PR-39 peptide used
13	90	100.0	39	22	AAW75722	Amino acid sequenc
14	90	100.0	39	22	AAW75722	PR-39 peptide. Un
15	90	100.0	42	23	AAW75722	Antimicrobial pept
16	90	100.0	44	22	AAW75722	E. coli AMP gene p
17	83	92.2	14	17	AAW75722	Leukocyte O2- prod
18	83	92.2	14	19	AAW75722	proline/Arginine r
19	75	83.3	23	17	AAW75722	Leukocyte O2- prod
20	66	73.3	18	16	AAW75722	Bactenecin peptide
21	66	73.3	20	19	AAW75722	Proline/Arginine r
22	66	73.3	23	16	AAW75722	Bactenecin peptide
23	66	73.3	35	16	AAW75722	Bactenecin peptide
24	66	73.3	59	19	AAW75722	Cationic peptide B
25	66	73.3	59	21	AAW75722	Cationic peptide B
26	66	73.3	60	23	AAW75722	Antimicrobial pept
27	66	73.3	62	22	AAW75722	Antimicrobial pept
28	64	71.1	11	21	AAW75722	E. coli AMP gene B
29	64	71.1	11	22	AAW75722	PR-39 derived angi
30	64	71.1	11	22	AAW75722	Amino acid sequenc
31	63	70.0	91	22	AAW75722	PR-39 derived pept
32	61	67.8	336	17	AAW75722	Propionibacterium
33	60	66.7	953	23	AAW75722	HCWV Toledo strain
34	60	66.7	953	23	AAW75722	Human protease PKT
35	59	65.6	39	21	AAW75722	Amino acid sequenc
36	59	65.6	59	17	AAW75722	Human secreted pro
37	56.5	62.8	74	22	AAW75722	Synducin peptide (
38	56.5	62.8	74	22	AAW75722	Peptide #10125 enc
39	56.5	62.8	74	22	AAW75722	Human brain expres
40	56.5	62.8	74	22	AAW75722	Human bone marrow
41	56.5	62.8	74	22	AAW75722	Peptide #10470 enc
42	56	62.2	692	22	AAW75722	Human peptide enco
43	55.5	61.7	497	20	AAW75722	Micromonospora eve
44	55	61.1	45	22	AAW75722	Mycobacterium spec
45	54	60.0	87	22	AAW75722	Peptide #5055 enco
						Propionibacterium

ALIGNMENTS

RESULT 1
AAW75722 standard; peptide; 15 AA.
ID AAW75722 standard; peptide; 15 AA.
XX AAW75722 standard; peptide; 15 AA.
AC AAW75722 standard; peptide; 15 AA.
XX AAW75722 standard; peptide; 15 AA.
DT 01-FEB-2001 (first entry)
XX PR-39 derived angiogenesis regulatory peptide 1.
DE PR-39 derived angiogenesis regulatory peptide 1.
XX Angiogenesis; stimulation; PR-39; anoxia; myocardial infarction;
KW myocardial ischaemia; proteasome.
XX Synthetic.
OS Synthetic.
PN WO200057895-A1.
XX WO200057895-A1.
PD 05-OCT-2000.
XX 16-MAR-2000; 2000WO-US07050.
PF 16-MAR-2000; 2000WO-US07050.
XX 26-MAR-1999; 99US-0276868.
XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
PI Simons M, Gao Y;
XX WPI; 2000-628319/60.
XX Stimulating angiogenesis in situ, useful e.g. for treating anoxia and
PT infarction, by administering a PR-39 oligopeptide that regulates
PT enzymatic activity of proteasomes

PS Claim 12; Page 40; 51pp; English.

CC This invention relates to a method for the stimulation of angiogenesis in

CC situ within a targeted collection of viable cells. The method comprises

CC introducing, into the cytoplasm, at least 1 member of the PR-39

CC oligopeptide collective, which interacts with cytoplasmic proteasomes.

CC Part of the proteolytic activity of the proteasomes is selectively

CC altered so as to stimulate angiogenesis. The method is used to induce

CC angiogenesis in tissue that has suffered anoxia or infarction,

CC e.g. myocardial infarction or chronic myocardial ischaemia, and also to

CC study the mechanisms that control angiogenesis. The present sequence

CC represents a PR-39 derived peptide which interacts with the proteasome

CC and can be used in the method of the invention.

XX

SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 21; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRPP 15

Db 1 RRRPRPPYLP RPRPP 15

|||||

RESULT 2

AAB84691

ID AAB84591 standard; peptide; 15 AA.

XX

AC AAB84691;

XX

DT 17-SEP-2001 (first entry)

XX

DE Amino acid sequence of a PR-39 derived peptide (residues 1-15).

XX

KW PR-39; IkappaBalpha degradation; NFkappaB transcription factor;

KW myocardial infarction; chronic myocardial ischemia; heart disease;

KW anoxia.

XX

OS Unidentified.

XX

PN WO200147540-A1.

XX

PD 05-JUL-2001.

XX

PF 27-DEC-2000; 2000WO-US35293.

XX

PR 29-DEC-1999; 99US-0474967.

XX

PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX

PI Simons M, Gao Y;

XX

DR WPI; 2001-441690/47.

XX

PT Selective inhibition of IkappaBalpha degradation within targeted viable

PT cell collection, involves interacting PR-39 oligopeptide with

PT IkappaBalpha and proteasomes, and altering proteolytic activity of

PT proteasomes -

XX

PS Claim 11; Page 58; 69pp; English.

XX

CC The present sequence represents a PR-39 derived peptide. It is used

CC for selective inhibition of IkappaBalpha degradation within a targeted

CC cell collection in-situ. The method is useful for selectively inhibiting

CC IkappaBalpha protein degradation in situ, decreasing the activity of

CC NFkappaB transcription factor and selective control of NFkappaB-dependent

CC gene expression in situ. The PR-39 derived peptides are useful in the

CC treatment of myocardial infarction, chronic myocardial ischemia of

CC heart disease and anoxia.

XX

SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRPP 15

Db 1 RRRPRPPYLP RPRPP 15

|||||

RESULT 3

AAB97277

ID AAB97277 standard; peptide; 15 AA.

XX

AC AAB97277;

XX

DT 09-AUG-2001 (first entry)

XX

DE PR-39 derived peptide PR-15.

XX

KW PR-39; cathelin; inflammation; wound healing; myocardial infarction;

KW proteasome; proteolysis; alpha7; peptide degradation; angiogenesis;

KW anoxia; chronic myocardial ischaemia; heart tissue.

XX

OS Unidentified.

XX

PN WO200130368-A1.

XX

PD 03-MAY-2001.

XX

PF 06-OCT-2000; 2000WO-US27552.

XX

PR 25-OCT-1999; 99US-0426011.

XX

PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX

PI Simons M, Gao Y;

XX

DR WPI; 2001-355179/37.

XX

PT Stimulation of angiogenesis and inhibition of proteasome mediated

PT degradation in cells, by introduction of PR-39 oligopeptide or its

PT N-terminal fragments or their conjugates, for use in anoxia and

PT infarction conditions -

XX

PS Claim 12; Page 42; 52pp; English.

XX

CC Peptides AAB97277 - AAB97279 represent PR-39 derived oligopeptides. PR-39

CC is a member of the the cathelin family of proteins, mature PR-39 is 39

CC amino acids in length (see AAB97280), and has been shown to play a role

CC in several inflammatory events including wound healing and myocardial

CC infarction. The PR-39 derived family of oligopeptides cause selective

CC inhibition of proteasome mediated degeneration of peptides and

CC stimulation of angiogenesis after their intracellular introduction to a

CC target cell. PR-39 derived peptides are able to interact with at least

CC the alpha7 subunit of the proteasomes, and therefore alter the

CC proteolytic activity of the proteasomes such that a selective increased

CC expression of specific proteins occurs. The invention includes methods

CC for the selective inhibition of proteasome mediated peptide degradation.

CC The method provides means for stimulating angiogenesis as required in

CC living tissues and organs which have suffered defects or have undergone

CC anoxia and/or infarction, myocardial infarction or chronic myocardial

CC ischaemia of heart tissue. Examples are the myocardium, skeletal or

CC smooth muscle, artery or vein, lung, brain, kidney, spleen, liver,

CC gastrointestinal or nerve tissues, limbs, and extremities. A particular

CC example is after myocardial infarction or ischaemia.

XX

SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRPP 15

Db 1 RRRPRPPLPRPRPP 15
 |||||

RESULT 4

AAW01452
 ID AAW01452 standard; peptide; 19 AA.

AC AAW01452;
 XX

DT 18-JUN-1997 (first entry)
 XX

DE Leukocyte O2- production inhibitor peptide PR19.
 XX

KW Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;
 KW antimicrobial peptide; small intestine; human; neutrophil; bacteria;
 KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;
 KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;
 KW tissue damage; oxygen radical; inflammatory disease; therapy.
 XX

OS Synthetic.
 XX

XX WO9632129-A1.
 PN

XX 17-OCT-1996.
 PD

XX 10-APR-1996; 96WO-US04674.
 PF

XX 10-APR-1995; 95US-0419066.
 PR

XX (UNIV) UNIV KANSAS STATE RES FOUND.
 PA

XX Blecha F, Shi J;
 PI

XX WPI; 1996-476842/47.
 DR

XX Inhibition of leukocyte super:oxide anion prodn. and attraction of
 PT leukocytes - using peptide(s) partic. based on antimicrobial PR-39
 PT

PS Disclosure; Page 27; 45pp; English.
 PS

XX AAW01447-W01454 represent fragments of the proline-arginine rich
 CC antimicrobial peptide PR39 (see AAW01446). The PR39 sequence was first
 CC isolated from porcine small intestine, and has also been identified in
 CC human and porcine neutrophils. PR39 kills bacteria by interfering with
 CC DNA and/or protein synthesis. PR39 also induces syndecan expression on
 CC mesenchymal cells. Syndecans are important in wound repair, showing that
 CC PR39 can be used in wound repair, as well as in antibacterial agents.
 CC These sequences, and PR39, can be used in the method of the invention.
 CC The method of the invention is for inhibiting leukocyte superoxide anion
 CC (O2-) production. The method comprises administering to a leukocyte a
 CC peptide (such as this sequence) capable of inhibiting leukocyte O2-
 CC production. The peptides can be used as medicaments for fighting
 CC infection by attracting leukocytes to a wound site and restricting
 CC tissue damage at the wound site caused by excessive oxygen radicals
 CC produced by these leukocytes. They can also be used to develop products
 CC for treating inflammatory disease states.
 XX

SQ Sequence 19 AA;
 XX

Query Match 100.0%; Score 90; DB 17; Length 19;
 Best Local Similarity 100.0%; Pred. No. 0.00025;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 RRRPRPPLPRPRPP 15
 |||||

Db 1 RRRPRPPLPRPRPP 15
 |||||

RESULT 5

AAW01447
 ID AAW01447 standard; peptide; 26 AA.

XX

AC AAW01447;
 XX

DT 18-JUN-1997 (first entry)
 XX

DE Leukocyte O2- production inhibitor peptide PR26.
 XX

KW Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;
 KW antimicrobial peptide; small intestine; human; neutrophil; bacteria;
 KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;
 KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;
 KW tissue damage; oxygen radical; inflammatory disease; therapy.
 XX

OS Synthetic.
 XX

XX WO9632129-A1.
 PN

XX 17-OCT-1996.
 PD

XX 10-APR-1996; 96WO-US04674.
 PF

XX 10-APR-1995; 95US-0419066.
 PR

XX (UNIV) UNIV KANSAS STATE RES FOUND.
 PA

XX Blecha F, Shi J;
 PI

XX WPI; 1996-476842/47.
 DR

XX Inhibition of leukocyte super:oxide anion prodn. and attraction of
 PT leukocytes - using peptide(s) partic. based on antimicrobial PR-39
 PT

PS Claim 3; Page 26; 45pp; English.
 PS

XX AAW01447-W01454 represent fragments of the proline-arginine rich
 CC antimicrobial peptide PR39 (see AAW01446). The PR39 sequence was first
 CC isolated from porcine small intestine, and has also been identified in
 CC human and porcine neutrophils. PR39 kills bacteria by interfering with
 CC DNA and/or protein synthesis. PR39 also induces syndecan expression on
 CC mesenchymal cells. Syndecans are important in wound repair, showing that
 CC PR39 can be used in wound repair, as well as in antibacterial agents.
 CC These sequences, and PR39, can be used in the method of the invention.
 CC The method of the invention is for inhibiting leukocyte superoxide anion
 CC (O2-) production. The method comprises administering to a leukocyte a
 CC peptide (such as this sequence) capable of inhibiting leukocyte O2-
 CC production. The peptides can be used as medicaments for fighting
 CC infection by attracting leukocytes to a wound site and restricting
 CC tissue damage at the wound site caused by excessive oxygen radicals
 CC produced by these leukocytes. They can also be used to develop products
 CC for treating inflammatory disease states.
 XX

SQ Sequence 26 AA;
 XX

Query Match 100.0%; Score 90; DB 17; Length 26;
 Best Local Similarity 100.0%; Pred. No. 0.00033;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 RRRPRPPLPRPRPP 15
 |||||

Db 1 RRRPRPPLPRPRPP 15
 |||||

RESULT 6

AAW75723
 ID AAW75723 standard; peptide; 26 AA.

XX AAW75723;
 AC

XX 19-NOV-1998 (first entry)
 DT

XX Proline/Arginine rich peptide PR-26.
 DE

XX Proline; arginine; peptide; reperfusion injury; neutrophil; endothelium;
 KW superoxide; neutrophil superoxide anion; chemotaxis; NADPH oxidase;
 KW

KW coronary bypass; organ transplantation surgery.
 XX Synthetic.
 OS
 XX WO9835690-A1.
 PN
 XX
 XX 20-AUG-1998.
 PD
 XX
 XX 17-FEB-1998; 98WO-US03207.
 PF
 XX
 XX 16-FEB-1998; 98US-0024975.
 PR
 XX 18-FEB-1997; 97US-0802306.
 PR
 XX (UNIV) UNIV KANSAS STATE RES FOUND.
 PA
 XX
 XX Blecha F, Ross CR, Shi J;
 PI
 XX WPI; 1998-495359/42.
 DR
 XX
 XX Reduction of reperfusion injury in temporarily occluded blood
 PT vessels - by administration of a peptide which is rich in proline
 PT or arginine residues
 PT
 XX
 PS Claim 3; Page 14-15; 35pp; English.
 XX
 CC Sequences AAW75722-W75732 are proline/arginine rich peptides that upon
 CC administration into a mammal's bloodstream reduce reperfusion injury
 CC (production of reactive oxygen species, neutrophil adherence to
 CC endothelium, and extravasation of neutrophils). These peptides have two
 CC requirements: they contain the consensus sequence PXXP, where P is a
 CC proline residue and X is any amino acid residue, which has been found to
 CC inhibit superoxide production, and secondly they have arginine residues
 CC adjacent to these motifs, required for effective inhibition. It was
 CC established by structural and function analysis that a peptide should
 CC ideally contain 4 or 6 of these motifs, and that inhibitory activity is
 CC correlated with the increase of length of peptides. The effectiveness
 CC of these peptides was determined by investigating the production of the
 CC neutrophil superoxide anion, and also the inhibition of neutrophil
 CC chemotaxis. From this, it was found that all of the peptides inhibited
 CC NADPH oxidase to some extent. All of the peptides also inhibit
 CC neutrophil oxidase activity. PR-39 is believed, to be the most potent
 CC endogenous down regulator of NADPH oxidase yet discovered, and from the
 CC data produced, it can be suggested to be involved in eliminating or
 CC reducing the reperfusion injury induced adhesion and extraction of
 CC neutrophils. The peptides are also useful in connection with surgical
 CC procedures such as coronary bypass and organ transplantation surgery.
 XX
 SQ Sequence 26 AA;
 Query Match 100.0%; Score 90; DB 19; Length 26;
 Best Local Similarity 100.0%; Pred. No. 0.00033;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLP RPP 15
 |||||
 Db 1 RRRPRPPYLP RPP 15
 RESULT 7
 AAR30491
 ID AAR30491 standard; peptide; 39 AA.
 XX
 XX AAR30491;
 AC
 XX
 XX 12-MAY-1993 (first entry)
 DT
 XX Antibacterial peptide.
 DE
 XX Pig; small intestine; endocrine; gram negative; bacteria; therapeutic;
 KW veterinary medicine; prophylactic.
 KW
 XX Sus scrofa domestica.
 OS
 XX

PN WO9222578-A.
 XX
 PD 23-DEC-1992.
 XX
 XX 10-JUN-1992; 92WO-SE00394.
 PF
 XX
 XX 14-JUN-1991; 91SE-0001838.
 PR
 XX (BOMA/) BOMAN H G.
 PA (JOER/) JOERNVALL H.
 PA (LEEJ/) LEE J.
 PA (MUTT/) MUTT V.
 XX
 XX Boman HG, Joernvall H, Lee J, Mutt V;
 PI
 XX WPI; 1993-018080/02.
 DR
 XX
 XX New anti-bacterial polypeptide - active against Gram negative
 PT bacteria
 PT
 XX
 PS Claim 1; Page 10; 15pp; English.
 XX
 CC This peptide was isolated from the small intestine of a pig. The
 CC small intestine is an important endocrine organ and many
 CC physiologically active peptides have been isolated from it. This
 CC peptide inhibits the growth of, and may kill, bacteria, pref. gram
 CC negative bacteria. This peptide or its functional derivatives may be
 CC used in human or veterinary medicine for therapeutic or prophylactic
 CC use.
 CC
 XX Sequence 39 AA;
 SQ
 Query Match 100.0%; Score 90; DB 14; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLP RPP 15
 |||||
 Db 1 RRRPRPPYLP RPP 15
 RESULT 8
 AAW01446
 ID AAW01446 standard; peptide; 39 AA.
 XX
 XX AAW01446;
 AC
 XX
 XX 18-JUN-1997 (first entry)
 DT
 XX
 XX Leukocyte O2- production inhibitor peptide PR39.
 DE
 XX
 XX Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;
 KW antimicrobial peptide; small intestine; human; neutrophil; bacteria;
 KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;
 KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;
 KW tissue damage; oxygen radical; inflammatory disease; therapy.
 XX
 OS Synthetic.
 OS
 XX
 XX WO9632129-A1.
 PN
 XX
 PD 17-OCT-1996.
 XX
 XX 10-APR-1996; 96WO-US04674.
 PF
 XX 10-APR-1995; 95US-0419066.
 PR
 XX (UNIV) UNIV KANSAS STATE RES FOUND.
 PA
 XX Blecha F, Shi J;
 PI
 XX WPI; 1996-476842/47.
 XX

PT Inhibition of leukocyte superoxide anion prodn. and attraction of
 PT leukocytes - using peptide(s) partic. based on antimicrobial PR-39
 PS
 PS Claim 2; Page 26; 45pp; English.
 XX
 CC This sequence represents the proline-arginine rich antimicrobial peptide
 CC PR39. The PR39 sequence was first isolated from porcine small intestine,
 CC and has also been identified in human and porcine neutrophils. PR39
 CC kills bacteria by interfering with DNA and/or protein synthesis. PR39
 CC also induces syndecan expression on mesenchymal cells. Syndecans are
 CC important in wound repair, showing that PR39 can be used in wound repair,
 CC as well as in antibacterial agents. This sequence, and the fragments of
 CC it shown in AA001447-W01454, can be used in the method of the invention.
 CC The method of the invention is for inhibiting leukocyte superoxide anion
 CC (O2-) production. The method comprises administering to a leukocyte a
 CC peptide (such as this sequence) capable of inhibiting leukocyte O2-
 CC production. The peptides can be used as medicaments for fighting
 CC infection by attracting leukocytes to a wound site and restricting tissue
 CC damage at the wound site caused by excessive oxygen radicals produced by
 CC these leukocytes. They can also be used to develop products for treating
 CC inflammatory disease states.
 XX
 SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 17; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPP 15
 |||||
 Db 1 RRRPRPPYLP RPP 15

RESULT 9

AA094446
 ID AAR94446 standard; peptide; 39 AA.

AC AAR94446;

DT 05-NOV-1996 (first entry)

DE Synducin peptide (PR-39) induces syndecan expression.

XX Synducin; induction; expression; syndecan-1; syndecan-4; surface;
 KW mesenchymal cell; fibroblast; epithelial; PR-39; treatment; stasis;
 KW decubitus; ulcers; keloids; skin burns; ischemic tissues;
 KW hypercoagulation states; prevention; tumour metastasis; restenosis;
 KW inhibition; angiogenesis; proliferation; endothelial.

OS Synthetic.

PN WO9609322-A2.

PD 28-MAR-1996.

PF 22-SEP-1995; 95WO-US12080.

PR 22-SEP-1994; 94US-0310722.

PA (CHIL-) CHILDRENS MEDICAL CENT.

PI Bernfield M, Gallo RL;

DR WPI; 1996-188401/19.

XX Modulating mesenchymal interaction by administration of synducan -
 PT used in the treatment of wounds, tumours, restenosis, etc

PS Claim 4; Page 26; 34pp; English.

XX The present peptide is a synducan, which induces the expression of
 CC syndecan-1 and syndecan-4 on the surface of mesenchymal cells, esp.
 CC fibroblasts and epithelial cells. The 36 N-terminal amino acids of

CC the peptide were found to be identical to the 36 N-terminal amino
 CC acids of PR-39, a Pro and Arg rich antibacterial peptide previously
 CC found in porcine intestine (WO9222578). Synducins may be used in
 CC the treatment of stasis and decubitus ulcers, keloids, skin burns,
 CC ischemic tissues and hypercoagulation states, prevention of tumour
 CC metastasis, restenosis inhibition and endothelial cell angiogenesis
 CC and proliferation induction.
 CC Human microvascular endothelial cells were assayed for syndecan-4
 CC expression following exposure to 5 % wound fluid, dbcAMP (1 mM),
 CC the present peptide (10 microm) or a blank, to give respective
 CC cell surface syndecan-4 values (MOD/m in) of approx. 1.75, 1.70,
 CC 1.80 and 0.95.

SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 17; Length 39;

Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPP 15
 |||||
 Db 1 RRRPRPPYLP RPP 15

RESULT 10

AA099121
 ID AAR99121 standard; peptide; 39 AA.

AC AAR99121;

DT 28-OCT-1996 (first entry)

DE Magainin-derived antimicrobial STD-inhibiting peptide, MSI-1312.

XX STD; sexually transmitted disease; HIV; human immunodeficiency virus;
 KW herpes simplex virus; HSV; Neisseria gonorrhoeae; Candida; Chlamydia;
 KW magainin; antimicrobial; squalamine.

OS Synthetic.

PH Key Location/Qualifiers
 FT Modified-site 39
 FT /note= "amidated"

PN WO9608270-A2.

PD 21-MAR-1996.

PF 13-SEP-1995; 95WO-US11675.

PR 13-SEP-1994; 94US-0305475.

PA (MAGA-) MAGAININ PHARM INC.

PI Bedi G, Jacob L, Williams T, Zasloff M;

DR WPI; 1996-179725/18.

XX Inhibiting sexually transmitted disease e.g. HIV or herpes simplex -
 PT by administering magainin antimicrobial or squalamine cpd. to
 PT inhibit transmission

PS Example 1; Page 32; 60pp; English.

XX AAR99116-R99123 are antimicrobial, magainin-analogue peptides that may
 CC be used to treat sexually transmitted diseases (STDs) caused by
 CC Chlamydia, HIV, herpes simplex virus, Neisseria gonorrhoeae or
 CC Candida infection. The peptides inhibit STDs by either killing the
 CC infectious organism, impeding the infection mechanism or
 CC interrupting the replication cycle of the organism. Squalamine (an
 CC aminosterol host defence molecule of the dog fish shark Squalus
 CC acanthias) and PGla (a frog antimicrobial peptide) analogues may
 CC also be useful in inhibiting STD infection and transmission.

SQ Sequence 39 AA;
 Query Match 100.0%; Score 90; DB 17; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
 |||||
 DB 1 RRRPRPPYLPRPRPP 15

RESULT 11
 AAW75722
 ID AAW75722 standard; peptide; 39 AA.
 XX
 AC AAW75722;
 XX
 DT 19-NOV-1998 (first entry)
 XX
 DE Proline/Arginine rich peptide PR-39.
 XX
 KW Proline; arginine; peptide; reperfusion injury; neutrophil; endothelium;
 KW superoxide; neutrophil superoxide anion; chemotaxis; NADPH oxidase;
 KW coronary bypass; organ transplantation surgery.
 XX
 OS Synthetic.
 XX
 PN WO9835690-A1.
 XX
 PD 20-AUG-1998.
 XX
 PF 17-FEB-1998; 98NO-US03207.
 XX
 PR 16-FEB-1998; 98US-0024975.
 PR 18-FEB-1997; 97US-0802306.
 XX
 PA (UNIV) UNIV KANSAS STATE RES FOUND.
 XX
 PI Blecha F, Ross CR, Shi J;
 XX
 DR WPI; 1998-495359/42.
 XX
 PT Reduction of reperfusion injury in temporarily occluded blood
 PT vessels - by administration of a peptide which is rich in proline
 PT or arginine residues
 XX
 Claim 3; Page 14; 35pp; English.

Sequences AAW75722-W75732 are proline/arginine rich peptides that upon
 administration into a mammal's bloodstream reduce reperfusion injury
 (production of reactive oxygen species, neutrophil adherence to
 endothelium, and extravasation of neutrophils). These peptides have two
 requirements: they contain the consensus sequence PXXP, where P is a
 proline residue and X is any amino acid residue, which has been found to
 inhibit superoxide production, and secondly they have arginine residues
 adjacent to these motifs, required for effective inhibition. It was
 established by structural and function analysis that a peptide should
 ideally contain 4 or 6 of these motifs, and that inhibitory activity is
 correlated with the increase of length of peptides. The effectiveness
 of these peptides was determined by investigating the production of the
 neutrophil superoxide anion, and also the inhibition of neutrophil
 chemotaxis. From this, it was found that all of the peptides inhibited
 NADPH oxidase to some extent. All of the peptides also inhibit
 neutrophil oxidase activity. PR-39 is believed, to be the most potent
 endogenous down regulator of NADPH oxidase yet discovered, and from the
 data produced, it can be suggested to be involved in eliminating or
 reducing the reperfusion injury induced adhesion and extraction of
 neutrophils. The peptides are also useful in connection with surgical
 procedures such as coronary bypass and organ transplantation surgery.

SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 19; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
 |||||
 DB 1 RRRPRPPYLPRPRPP 15

RESULT 12
 AAB26888
 ID AAB26888 standard; peptide; 39 AA.
 XX
 AC AAB26888;
 XX
 DT 01-FEB-2001 (first entry)
 XX
 DE PR-39 peptide used in angiogenesis control.
 XX
 KW Angiogenesis; stimulation; PR-39; anoxia; myocardial infarction;
 KW myocardial ischaemia; proteasome.
 XX
 OS Synthetic.
 XX
 PN WO200057895-A1.
 XX
 PD 05-OCT-2000.
 XX
 PF 16-MAR-2000; 2000WO-US07050.
 XX
 PR 26-MAR-1999; 99US-0276868.
 XX
 PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
 XX
 PI Simons M, Gao Y;
 XX
 DR WPI; 2000-628319/60.
 XX
 PT Stimulating angiogenesis in situ, useful e.g. for treating anoxia and
 PT infarction, by administering a PR-39 oligopeptide that regulates
 PT enzymatic activity of proteasomes
 XX
 PS Disclosure; Page 21; 51pp; English.

This invention relates to a method for the stimulation of angiogenesis in
 situ within a targeted collection of viable cells. The method comprises
 introducing, into the cytoplasm, at least 1 member of the PR-39
 oligopeptide collective, which interacts with cytoplasmic proteasomes.
 Part of the proteolytic activity of the proteasomes is selectively
 altered so as to stimulate angiogenesis. The method is used to induce
 angiogenesis in tissue that has suffered anoxia or infarction,
 e.g. myocardial infarction or chronic myocardial ischaemia, and also to
 study the mechanisms that control angiogenesis. The present sequence
 represents the PR-39 peptide from which peptide used in the method of
 the invention are derived.

SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 21; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
 |||||
 DB 1 RRRPRPPYLPRPRPP 15

RESULT 13
 AAB84690
 ID AAB84690 standard; protein; 39 AA.
 XX
 AC AAB84690;
 XX

DT 17-SEP-2001 (first entry)
 DE Amino acid sequence of a PR-39 protein.
 XX
 KW PR-39; IkappaBalpHa degradation; NFkappaB transcription factor;
 KW myocardial infarction; chronic myocardial ischemia; heart disease;
 KW anoxia.
 XX
 OS Unidentified.
 XX
 PN WO200147540-A1.
 XX
 PD 05-JUL-2001.
 XX
 PF 27-DEC-2000; 2000WO-US35293.
 XX
 PR 29-DEC-1999; 99US-0474967.
 XX
 PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
 XX
 PI Simons M, Gao Y;
 XX
 DR WPI; 2001-441690/47.
 XX
 XX Selective inhibition of IkappaBalpHa degradation within targeted viable
 PT cell collection, involves interacting PR-39 oligopeptide with
 PT IkappaBalpHa and proteasomes, and altering proteolytic activity of
 PT proteasomes.
 XX
 PS Disclosure; Page 30; 69pp; English.
 XX
 CC The present sequence represents a PR-39 protein. The specification
 CC describes PR-39 derived peptides, which are used for selective
 CC inhibition of IkappaBalpHa degradation within a targeted cell collection
 CC in-situ. The method is useful for selectively inhibiting IkappaBalpHa
 CC protein degradation in situ, decreasing the activity of NFkappaB
 CC transcription factor and selective control of NFkappaB-dependent gene
 CC expression in situ. The PR-39 derived peptides are useful in the
 CC treatment of myocardial infarction, chronic myocardial ischemia of
 CC heart disease and anoxia.
 XX
 SQ Sequence 39 AA;
 Query Match 100.0%; Score 90; DB 22; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLP RPP 15
 DB 1 RRRPRPPYLP RPP 15
 RESULT 14
 AAB97280
 ID AAB97280 standard; peptide; 39 AA.
 XX
 AC AAB97280;
 XX
 DT 09-AUG-2001 (first entry)
 XX
 DE PR-39 peptide.
 XX
 KW PR-39; cathelin; inflammation; wound healing; myocardial infarction;
 KW proteasome; proteolysis; alpha7; peptide degradation; angiogenesis;
 KW anoxia; chronic myocardial ischemia; heart tissue.
 XX
 OS Unidentified.
 XX
 PN WO200130368-A1.
 XX
 PD 03-MAY-2001.
 XX
 PF 06-OCT-2000; 2000WO-US27552.
 XX

XX 25-OCT-1999; 99US-0426011.
 XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
 XX
 PI Simons M, Gao Y;
 XX
 DR WPI; 2001-355179/37.
 XX
 XX Stimulation of angiogenesis and inhibition of proteasome mediated
 PT degradation in cells, by introduction of PR-39 oligopeptide or its
 PT N-terminal fragments or their conjugates, for use in anoxia and
 PT infarction conditions.
 XX
 PS Disclosure; Page 21; 52pp; English.
 XX
 CC Peptides AAB97277 - AAB97279 represent PR-39 derived oligopeptides. PR-39
 CC is a member of the the cathelin family of proteins, mature PR-39
 CC represented by the present sequence is 39 amino acids in length, and has
 CC been shown to play a role in several inflammatory events including wound
 CC healing and myocardial infarction. The PR-39 derived family of
 CC oligopeptides cause selective inhibition of proteasome mediated
 CC degeneration of peptides and stimulation of angiogenesis after their
 CC intracellular introduction to a target cell. PR-39 derived peptides are
 CC able to interact with at least the alpha7 subunit of the proteasomes, and
 CC therefore alter the proteolytic activity of proteasomes such that a
 CC selective increased expression of specific proteins occurs. The invention
 CC includes methods for the selective inhibition of proteasome mediated
 CC peptide degradation. The method provides means for stimulating
 CC angiogenesis as required in living tissues and organs which have suffered
 CC defects or have undergone anoxia and/or infarction, myocardial infarction
 CC or chronic myocardial ischemia of heart tissue. Examples are the
 CC myocardium, skeletal or smooth muscle, artery or vein, lung, brain,
 CC kidney, spleen, liver, gastrointestinal or nerve tissues, limbs, and
 CC extremities. A particular example is after myocardial infarction or
 CC ischaemia.
 XX
 SQ Sequence 39 AA;
 Query Match 100.0%; Score 90; DB 22; Length 39;
 Best Local Similarity 100.0%; Pred. No. 0.00047;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLP RPP 15
 DB 1 RRRPRPPYLP RPP 15
 RESULT 15
 ABB07714
 ID ABB07714 standard; peptide; 42 AA.
 XX
 AC ABB07714;
 XX
 DT 10-JUN-2002 (first entry)
 XX
 DE Antimicrobial peptide PR-39 C-terminal fragment.
 XX
 KW Vaccine; cathelicidin; antimicrobial; immunostimulant; immune response;
 KW antigen presenting cell; adjuvant; porcine; PR-39.
 XX
 OS Sus sp.
 XX
 PN WO200213857-A2.
 XX
 PD 21-FEB-2002.
 XX
 PF 17-AUG-2001; 2001WO-EP09529.
 XX
 PR 17-AUG-2000; 2000AT-0001416.
 XX
 PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX

PI Fritz J, Mattner F, Zauner W, Buschle M, Egyed A;
 XX
 DR WPI; 2002-269154/31.
 XX
 PT Vaccine for active immunization or for preparing an adjuvant for
 PT enhancing an immune response to at least one antigen, comprises at
 PT least one antigen and at least one cathelicidin derived antimicrobial
 PT peptide -
 XX
 PS Disclosure; Fig 3; 65pp; English.
 XX
 CC The invention relates to a vaccine comprising at least one antigen and at
 CC least one cathelicidin derived antimicrobial peptide or its derivative.
 CC The vaccine is useful for active immunization, especially of humans or
 CC animals without protection against the specific antigen. The cathelicidin
 CC derived antimicrobial peptide is useful in the preparation of an adjuvant
 CC for enhancing the immune response to at least one antigen, where the
 CC adjuvant enhances the uptake of at least one antigen in antigen
 CC presenting cells (APC), and the adjuvant is added to the vaccine.
 CC Sequences ABB07708-15 represent C-terminal fragments of antimicrobial
 CC peptides of the cathelicidin family.
 XX
 SQ Sequence 42 AA;
 Query Match 100.0%; Score 90; DB 23; Length 42;
 Best Local Similarity 100.0%; Pred. No. 0.0005;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLP RPP 15
 Db 1 RRRPRPPYLP RPP 15
 |||||
 |||||

Search completed: May 13, 2003, 10:40:32
 Job time : 36 secs

GenCore version 5.1.4 p5 4578
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:40:17 ; Search time 14 Seconds
(without alignments)
31.525 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPPYLPRLPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 262574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents AA:*

- 1: /cgn2_6/prodata/1/iaa/5A_COMB.pep:*
- 2: /cgn2_6/prodata/1/iaa/5B_COMB.pep:*
- 3: /cgn2_6/prodata/1/iaa/6A_COMB.pep:*
- 4: /cgn2_6/prodata/1/iaa/6B_COMB.pep:*
- 5: /cgn2_6/prodata/1/iaa/6C_COMB.pep:*
- 6: /cgn2_6/prodata/1/iaa/backfiles1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	90	100.0	26	US-08-419-066-2	Sequence 2, Appli
2	90	100.0	26	US-09-024-975-2	Sequence 2, Appli
3	90	100.0	39	US-08-162-052-1	Sequence 1, Appli
4	90	100.0	39	US-08-310-722-1	Sequence 1, Appli
5	90	100.0	39	US-08-419-066-1	Sequence 1, Appli
6	90	100.0	39	US-08-728-333-1	Sequence 1, Appli
7	90	100.0	39	US-09-024-975-1	Sequence 1, Appli
8	90	100.0	39	PCT-US95-12080-1	Sequence 1, Appli
9	83	92.2	14	US-09-024-975-4	Sequence 4, Appli
10	66	73.3	20	US-09-024-975-9	Sequence 9, Appli
11	61	67.8	336	US-08-414-926A-26	Sequence 26, Appl
12	61	67.8	336	US-08-926-922-26	Sequence 26, Appl
13	61	67.8	336	US-09-253-682-26	Sequence 26, Appl
14	61	67.8	336	US-09-527-657-26	Sequence 26, Appl
15	59	65.6	59	PCT-US95-12080-3	Sequence 3, Appli
16	53	58.9	18	US-08-205-938A-23	Sequence 23, Appl
17	53	58.9	18	US-08-205-938A-24	Sequence 24, Appl
18	53	58.9	18	US-09-230-180-20	Sequence 20, Appl
19	53	58.9	18	PCT-US95-02626-23	Sequence 23, Appl
20	53	58.9	18	PCT-US95-02626-24	Sequence 24, Appl
21	52	57.8	18	US-08-205-938A-25	Sequence 25, Appl
22	52	57.8	18	PCT-US95-02626-25	Sequence 25, Appl
23	51.5	57.2	355	US-08-483-533-41	Sequence 41, Appl
24	51.5	57.2	355	US-09-283-471A-41	Sequence 41, Appl
25	51.5	57.2	355	PCT-US91-06532-3	Sequence 3, Appli
26	51	56.7	16	US-08-205-938A-8	Sequence 8, Appli
27	51	56.7	16	PCT-US95-02626-8	Sequence 8, Appli

28	51	56.7	180	3	US-09-187-331-5	Sequence 5, Appli
29	51	56.7	180	4	US-09-470-946-5	Sequence 5, Appli
30	51	56.7	195	3	US-09-187-331-1	Sequence 1, Appli
31	51	56.7	195	4	US-09-470-946-1	Sequence 1, Appli
32	50.5	56.1	169	4	US-08-483-533-28	Sequence 28, Appl
33	50.5	56.1	169	4	US-09-283-471A-28	Sequence 28, Appl
34	50.5	56.1	393	4	US-09-432-470-2	Sequence 2, Appli
35	50.5	56.1	393	4	US-09-432-470-4	Sequence 4, Appli
36	50	55.6	16	1	US-08-205-938A-7	Sequence 7, Appli
37	50	55.6	16	1	US-08-205-938A-28	Sequence 28, Appl
38	50	55.6	16	5	PCT-US95-02626-7	Sequence 7, Appli
39	50	55.6	16	5	PCT-US95-02626-28	Sequence 28, Appl
40	50	55.6	17	1	US-08-205-938A-27	Sequence 27, Appl
41	50	55.6	17	5	PCT-US95-02626-27	Sequence 27, Appl
42	49.5	55.0	129	4	US-09-199-637A-97	Sequence 97, Appl
43	49	54.4	26	4	US-09-024-975-8	Sequence 8, Appli
44	48.5	53.9	716	4	US-09-186-276B-67	Sequence 67, Appl
45	48.5	53.9	716	4	US-08-842-445-67	Sequence 67, Appl

ALIGNMENTS

RESULT 1
US-08-419-066-2
; Sequence 2, Application US/08419066
; Patent No, 5830993
; GENERAL INFORMATION:
; APPLICANT: Blecha, Frank
; APPLICANT: Shi, Jishu
; TITLE OF INVENTION: SYNTHETIC ANTIMICROBIAL PEPTIDE
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John M. Collins, Hovey, Williams, Timmons &
; ADDRESSEE: Collins
; STREET: 2405 Grand Boulevard, Suite 400
; CITY: Kansas City
; STATE: Missouri
; COUNTRY: U.S.A.
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/419,066
; FILING DATE:
; CLASSIFICATION: 530
; ATTORNEY/AGENT INFORMATION:
; NAME: Collins, John M.
; REGISTRATION NUMBER: 26262
; REFERENCE/DOCKET NUMBER: 23625
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (816) 474-9050
; TELEFAX: (816) 474-9057
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 26 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: N-terminal
US-08-419-066-2

Query Match 100.0%; Score 90; DB 2; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.2e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 RRRPPYLPRLPP 15

Db 1 RRRPPYLPYLPYLP 15

RESULT 2

US-09-024-975-2
; Sequence 2, Application US/09024975
; Patent No. 6133233
; GENERAL INFORMATION:
; APPLICANT: ROSS, CHRISTOPHER R.
; APPLICANT: BLECHA, FRANK
; APPLICANT: SHI, JISHU
; TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
; STREET: 2405 GRAND BLVD., SUITE 400
; CITY: KANSAS CITY
; STATE: MO
; COUNTRY: USA
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/024,975
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/802,306
; FILING DATE: 18-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: COLLINS, JOHN M.
; REGISTRATION NUMBER: 26,262
; REFERENCE/DOCKET NUMBER: 25585-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 816/474-9050
; TELEFAX: 816/474-9057
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 26 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-024-975-2

Query Match 100.0%; Score 90; DB 4; Length 26;
Best Local Similarity 100.0%; Pred. No. 6.2e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 RRRPPYLPYLPYLP 15
Db 1 RRRPPYLPYLPYLP 15

RESULT 3

US-08-162-052-1
; Sequence 1, Application US/08162052
; Patent No. 5489575
; GENERAL INFORMATION:
; APPLICANT: LEE, Jong-Youn
; APPLICANT: BOWAN, Hans G
; APPLICANT: MUTT, Viktor
; APPLICANT: JORNVAL, Hans
; TITLE OF INVENTION: NOVEL POLYPEPTIDES AND THEIR USE
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Burns, Doane, Swecker & Mathis
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia

COUNTRY: United States
ZIP: 22313-1404
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/162,052
FILING DATE: 02-JUN-1994
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: SE 9101838-2
FILING DATE: 14-JUN-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO 92-22578
FILING DATE: 23-DEC-1992
ATTORNEY/AGENT INFORMATION:
NAME: Crane-Feury, Sharon E
REGISTRATION NUMBER: 36,113
REFERENCE/DOCKET NUMBER: 003300-299
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 836-6620
TELEFAX: (703) 836-2021
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-162-052-1

Query Match 100.0%; Score 90; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 RRRPPYLPYLPYLP 15
Db 1 RRRPPYLPYLPYLP 15

RESULT 4

US-08-310-722-1
; Sequence 1, Application US/08310722
; Patent No. 5654273
; GENERAL INFORMATION:
; APPLICANT: Gallo, Richard L.
; APPLICANT: Klagsbrun, Michael
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabst
; STREET: 1100 Peachtree Street, Suite 2800
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-4530
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/310,722
; FILING DATE: 22-SEP-1994
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
; REFERENCE/DOCKET NUMBER: CMCC379
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-815-6508

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; TELEFAX: (404)-815-6555
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PUBLICATION INFORMATION:
; AUTHORS: Lee, Jong-Youn
; AUTHORS: Boman, Hans G.
; AUTHORS: Mutt, Viktor
; AUTHORS: Jornvall, Hans
; TITLE: No. 5654273el Polypeptides And Their Use
; JOURNAL: PCT WO 92/22578
; DATE: 12/23/92
; RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
; US-08-310-722-1

Query Match 100.0%; Score 90; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPPYLPYLRPP 15
   |||||
Db 1 RRRPPYLPYLRPP 15

RESULT 5
US-08-419-066-1
; Sequence 1, Application US/08419066
; Patent No. 5830993
; GENERAL INFORMATION:
; APPLICANT: Blecha, Frank
; APPLICANT: Shi, Jishu
; TITLE OF INVENTION: SYNTHETIC ANTIMICROBIAL PEPTIDE
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John M. Collins, Hovey, Williams, Timmons &
; ADDRESSEE: Collins
; STREET: 2405 Grand Boulevard, Suite 400
; CITY: Kansas City
; STATE: Missouri
; COUNTRY: U.S.A.
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/419,066
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Collins, John M.
; REGISTRATION NUMBER: 26262
; REFERENCE/DOCKET NUMBER: 23625
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (816) 474-9050
; TELEFAX: (816) 474-9057
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: N-terminal
; US-08-419-066-1

Query Match 100.0%; Score 90; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPPYLPYLRPP 15
   |||||
Db 1 RRRPPYLPYLRPP 15

RESULT 6
US-08-728-333-1
; Sequence 1, Application US/08728333
; Patent No. 5863897
; GENERAL INFORMATION:
; APPLICANT: Gallo, Richard L.
; APPLICANT: Klagsbrun, Michael
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabet
; STREET: 1100 Peachtree Street, Suite 2800
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-4530
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/728,333
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/310,722
; FILING DATE: 22-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
; REFERENCE/DOCKET NUMBER: CMCC379
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-815-6508
; TELEFAX: (404)-815-6555
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PUBLICATION INFORMATION:
; AUTHORS: Lee, Jong-Youn
; AUTHORS: Boman, Hans G.
; AUTHORS: Mutt, Viktor
; AUTHORS: Jornvall, Hans
; TITLE: No. 5863897el Polypeptides And Their Use
; JOURNAL: PCT WO 92/22578
; DATE: 12/23/92
; RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
; US-08-728-333-1

Query Match 100.0%; Score 90; DB 2; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPPYLPYLRPP 15
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Db 1 RRRPPYLPYLRPP 15
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RESULT 7
US-09-024-975-1
; Sequence 1, Application US/09024975
; Patent No. 6133233
; GENERAL INFORMATION:
; APPLICANT: ROSS, CHRISTOPHER R.
; APPLICANT: BLECHA, FRANK
; APPLICANT: SHI, JISHU
; TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
; STREET: 2405 GRAND BLVD., SUITE 400
; CITY: KANSAS CITY
; STATE: MO
; COUNTRY: USA
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/024,975
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/802,306
; FILING DATE: 18-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: COLLINS, JOHN M.
; REGISTRATION NUMBER: 26,262
; REFERENCE/DOCKET NUMBER: 25585-A
; TELEPHONE: 816/474-9050
; TELEFAX: 816/474-9057
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-024-975-1
Query Match 100.0%; Score 90; DB 4; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 RRRPRPPYLP RPP 15
| | | | | | | | | |
DB 1 RRRPRPPYLP RPP 15
| | | | | | | | | |
RESULT 8
PCT-US95-12080-1
; Sequence 1, Application PC/TUS9512080
; GENERAL INFORMATION:
; APPLICANT: Children's Medical Center Corporation
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabst
; STREET: 2800 One Atlantic Center
; STREET: 1201 West Peachtree
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-3450
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
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; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/12080
; FILING DATE:
; CLASSIFICATION:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-873-8794
; TELEFAX: (404)-815-8795
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PUBLICATION INFORMATION:
; AUTHORS: Lee, Jong-Youn
; AUTHORS: Boman, Hans G.
; AUTHORS: Mutt, Viktor
; AUTHORS: Jornvall, Hans
; TITLE: Novel Polypeptides And Their Use
; JOURNAL: PCT WO 92/22578
; DATE: 12/23/92
; RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
PCT-US95-12080-1
Query Match 100.0%; Score 90; DB 5; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 RRRPRPPYLP RPP 15
| | | | | | | | | |
DB 1 RRRPRPPYLP RPP 15
| | | | | | | | | |
RESULT 9
US-09-024-975-4
; Sequence 4, Application US/09024975
; Patent No. 6133233
; GENERAL INFORMATION:
; APPLICANT: ROSS, CHRISTOPHER R.
; APPLICANT: BLECHA, FRANK
; APPLICANT: SHI, JISHU
; TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
; STREET: 2405 GRAND BLVD., SUITE 400
; CITY: KANSAS CITY
; STATE: MO
; COUNTRY: USA
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/024,975
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/802,306
; FILING DATE: 18-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: COLLINS, JOHN M.
; REGISTRATION NUMBER: 26,262
; REFERENCE/DOCKET NUMBER: 25585-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 816/474-9050
; TELEFAX: 816/474-9057
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
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LENGTH: 14 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-975-4

Query Match 92.2%; Score 83; DB 4; Length 14;
Best Local Similarity 100.0%; Pred. No. 0.00023;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPP 14
DB 1 RRRPRPPLPRPP 14

RESULT 10
US-09-024-975-9
Sequence 9, Application US/09024975
Patent No. 6133233
GENERAL INFORMATION:
APPLICANT: ROSS, CHRISTOPHER R.
APPLICANT: BLECHA, FRANK
TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
NUMBER OF SEQUENCES: 11
CORRESPONDENCE ADDRESS:
ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
STREET: 2405 GRAND BLVD., SUITE 400
CITY: KANSAS CITY
STATE: MO
COUNTRY: USA
ZIP: 64108
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,975
FILING DATE:

CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/802,306
FILING DATE: 18-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: COLLINS, JOHN M.
REGISTRATION NUMBER: 26,262
REFERENCE/DOCKET NUMBER: 25585-A
TELECOMMUNICATION INFORMATION:
TELEPHONE: 816/474-9050
TELEFAX: 816/474-9057
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-975-9

Query Match 73.3%; Score 66; DB 4; Length 20;
Best Local Similarity 85.7%; Pred. No. 0.032;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPP 14
DB 2 RRRPRPPLPRPP 15

RESULT 11
US-08-414-926A-26
Sequence 26, Application US/08414926A
Patent No. 5721354
GENERAL INFORMATION:

APPLICANT: Spaete, Richard
APPLICANT: Cha, Tai-An
TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS
NUMBER OF SEQUENCES: 27
CORRESPONDENCE ADDRESS:
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum
STREET: 5 Palo Alto Square
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94306-2155
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/414,926A
FILING DATE: March 31, 1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Cserr, Luann
REGISTRATION NUMBER: 31,822
REFERENCE/DOCKET NUMBER: AVIR-011/OOUS
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-494-7622
TELEFAX: 415-857-0663
INFORMATION FOR SEQ ID NO: 26:
SEQUENCE CHARACTERISTICS:
LENGTH: 336 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
IMMEDIATE SOURCE:
CLONE: tol.22
FEATURE:
NAME/KEY: Protein
LOCATION: 1..336
OTHER INFORMATION: /label= UL151
US-08-414-926A-26

Query Match 67.8%; Score 61; DB 1; Length 336;
Best Local Similarity 78.6%; Pred. No. 1.7;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 RRRPRPPLPRPP 15
DB 279 RRRPILQPRPP 292

RESULT 12
US-08-926-922-26
Sequence 26, Application US/08926922
Patent No. 5925751
GENERAL INFORMATION:
APPLICANT: Spaete, Richard
APPLICANT: Cha, Tai-An
TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS
NUMBER OF SEQUENCES: 27
CORRESPONDENCE ADDRESS:
ADDRESSEE: Luann Cserr Attorney at Law
STREET: 750 Arimo Avenue
CITY: Oakland
STATE: CA
COUNTRY: USA
ZIP: 94610
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/926,922

Db 279 RREIPFLOPRPP 292

RESULT 15

PCT-US95-12080-3
; Sequence 3, Application PC/TUS9512080
; GENERAL INFORMATION:
; APPLICANT: Children's Medical Center Corporation
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabst
; STREET: 2800 One Atlantic Center
; STREET: 1201 West Peachtree
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-3450
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/12080
; FILING DATE:
; CLASSIFICATION:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-873-8794
; TELEFAX: (404)-815-8795
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 59 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; PCT-US95-12080-3

Query Match 65.6%; Score 59; DB 5; Length 59;
Best Local Similarity 84.6%; Pred. No. 0.57;
Matches 11; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPFR 13
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Db 2 RIRPRPRLPRR 14

Search completed: May 13, 2003, 10:42:08
Job time : 15 secs

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GenCore version 5.1.4 p5 4578
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:40:37 ; Search time 17 seconds
(without alignments)
81.199 Million cell updates/sec

Title: US-09-426-011D-3

Perfect score: 90

Sequence: 1 RRRPRPPYLPRLPRP 15

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Gapop 10.0 , Gapext 0.5

Searched: 349150 seqs, 92025710 residues

Total number of hits satisfying chosen parameters: 349150

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Published Applications AA:*

- 1: /cgn2_6/ptodata/2/pubpaa/US08_NEW PUB.pap.*
- 2: /cgn2_6/ptodata/2/pubpaa/PCT_NEW PUB.pap.*
- 3: /cgn2_6/ptodata/2/pubpaa/US06_NEW PUB.pap.*
- 4: /cgn2_6/ptodata/2/pubpaa/US06_PUBCOMB.pap.*
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- 8: /cgn2_6/ptodata/2/pubpaa/US08_PUBCOMB.pap.*
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- 10: /cgn2_6/ptodata/2/pubpaa/US09_PUBCOMB.pap.*
- 11: /cgn2_6/ptodata/2/pubpaa/US10_NEW PUB.pap.*
- 12: /cgn2_6/ptodata/2/pubpaa/US10_PUBCOMB.pap.*
- 13: /cgn2_6/ptodata/2/pubpaa/US60_NEW PUB.pap.*
- 14: /cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	66	73.3	59	10	US-09-030-619-163
2	60	66.7	953	10	US-09-888-615-66
3	56.5	62.8	74	10	US-09-864-761-45555
4	55	61.1	45	10	US-09-864-761-49065
5	54	60.0	250	9	US-10-102-806-517
6	53	58.9	18	10	US-09-030-619-96
7	53	58.9	18	10	US-09-030-619-158
8	53	58.9	18	10	US-09-030-619-159
9	52	57.8	18	10	US-09-030-619-160
10	52	57.8	354	9	US-10-004-717-2
11	52	57.8	354	9	US-10-004-717-58
12	51	56.7	180	10	US-09-997-701-5
13	51	56.7	195	10	US-09-997-701-1
14	50.5	56.1	392	10	US-09-747-835A-55
15	50.5	56.1	393	9	US-10-243-035-2
16	50.5	56.1	419	10	US-09-828-035-2
17	50.5	56.1	1314	10	US-09-747-835A-29
18	50	55.6	99	10	US-09-864-761-43778
19	50	55.6	146	9	US-09-989-920-237

20	50	55.6	449	9	US-10-125-540-320	Sequence 320, App
21	50	55.6	449	9	US-10-103-313-438	Sequence 438, App
22	50	55.6	449	10	US-09-764-870-320	Sequence 320, App
23	50	55.6	449	10	US-09-764-853-643	Sequence 643, App
24	49.5	55.0	129	9	US-09-975-719-97	Sequence 97, Appl
25	49	54.4	223	12	US-10-062-254-204	Sequence 204, App
26	48.5	53.9	111	10	US-09-864-761-47005	Sequence 47005, A
27	48	53.3	281	8	US-08-971-317A-6	Sequence 6, Appli
28	48	53.3	281	9	US-09-131-237-6	Sequence 6, Appli
29	48	53.3	281	9	US-10-174-554-10	Sequence 10, Appl
30	48	53.3	281	9	US-10-151-882-44	Sequence 44, Appl
31	48	53.3	281	10	US-09-802-669-25	Sequence 25, Appl
32	48	53.3	281	10	US-09-193-663-6	Sequence 6, Appli
33	48	53.3	281	10	US-09-037-287-6	Sequence 6, Appli
34	48	53.3	281	10	US-09-252-656B-6	Sequence 6, Appli
35	48	53.3	281	10	US-09-929-493-6	Sequence 6, Appli
36	48	53.3	281	10	US-09-927-110-1	Sequence 1, Appli
37	48	53.3	281	12	US-10-012-452-13	Sequence 13, Appl
38	48	53.3	1134	9	US-10-001-873-50	Sequence 50, Appl
39	47.5	52.8	272	10	US-09-925-300-1697	Sequence 1697, Ap
40	47	52.2	42	10	US-09-030-619-162	Sequence 162, App
41	47	52.2	84	9	US-10-174-590-486	Sequence 486, App
42	47	52.2	84	9	US-10-176-758-486	Sequence 486, App
43	47	52.2	84	9	US-10-175-737-486	Sequence 486, App
44	47	52.2	84	9	US-10-173-706-486	Sequence 486, App
45	47	52.2	84	9	US-10-175-738-486	Sequence 486, App

ALIGNMENTS

RESULT 1

US-09-030-619-163

; Sequence 163, Application US/09030619B

; Patent No. US20020035061A1

; GENERAL INFORMATION:

; APPLICANT: Krieger, Timothy J.

; APPLICANT: Taylor, Robert

; APPLICANT: Erfile, Douglas

; APPLICANT: Fraser, Janet R.

; APPLICANT: West, Michael H.P.

; APPLICANT: McNicol, Patricia J.

; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING

; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION

; TITLE OF INVENTION: WITH ANTIBIOTICS

; FILE REFERENCE: 66081.406

; CURRENT APPLICATION NUMBER: US/09/030,619B

; CURRENT FILING DATE: 1998-02-25

; NUMBER OF SEQ ID NOS: 232

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 163

; LENGTH: 59

; TYPE: PRT

; ORGANISM: Bos taurus

; US-09-030-619-163

Query Match 73.3%; Score 66; DB 10; Length 59;

Best Local Similarity 85.7%; Pred. No. 0.27; Mismatches 2; Indels 0; Gaps 0;

Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 RRRPRPPYLPRLPRP 14

| | | | | | | | | |

Db 2 RRRPRPPYLPRLPRP 15

| | | | | | | | | |

RESULT 2

US-09-888-615-66

; Sequence 66, Application US/09888615

; Patent No. US20020064856A1

; GENERAL INFORMATION:

; APPLICANT: FLOWMAN, GREGORY

; APPLICANT: WHYTE, DAVID

; APPLICANT: CAENEPEEL, SEAN

APPLICANT: CHARYDCZAK, GLEN
APPLICANT: MANNING, GERARD
APPLICANT: SUDARSANAM, SUCHA
TITLE OF INVENTION: NOVEL PROTEASES
FILE REFERENCE: 038602/1214
CURRENT APPLICATION NUMBER: US/09/888,615
CURRENT FILING DATE: 2001-06-26
PRIOR APPLICATION NUMBER: 60/214,047
PRIOR FILING DATE: 2000-06-26
NUMBER OF SEQ ID NOS: 150
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 66
LENGTH: 953
TYPE: PRT
ORGANISM: Homo sapiens
US-09-888-615-66

Query Match 66.7%; Score 60; DB 10; Length 953;
Best Local Similarity 56.5%; Pred. No. 17;
Matches 13; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 RRRPRP-----PYLPRRPP 15
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Db 377 RRRPRPQTLRTPQPQRRPP 399

RESULT 3
US-09-864-761-45555
Sequence 45555, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharron G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeonica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60/234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408

PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29
NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
SEQ ID NO 45555
LENGTH: 74
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AC010458.2
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.1
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 0.9
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.2
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 0.79
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.2
OTHER INFORMATION: EST_HUMAN HIT: AW503858.1, EVALUATE 5.00e-20
US-09-864-761-45555

Query Match 62.8%; Score 56.5; DB 10; Length 74;
Best Local Similarity 56.2%; Pred. No. 3.9;
Matches 9; Conservative 4; Mismatches 2; Indels 1; Gaps 1;

QY 1 RRRPRPPY-LRRPRPP 15
||||| :|||: :|||:
Db 37 RRRPKPPHRIPEPKPP 52

RESULT 4
US-09-864-761-49065
Sequence 49065, Application US/09864761
Patent No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharron G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeonica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00670

; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 49065
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC005973.2
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.88
; OTHER INFORMATION: EST_HUMAN HIT: A1358103.1, EVALUATE 4.60e+00
US-09-864-761-49065

Query Match 61.1%; Score 55; DB 10; Length 45;
Best Local Similarity 76.9%; Pred. No. 3.6;
Matches 10; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPR 13
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Db 19 RRRPRPPGPRPP 31

RESULT 5

US-10-102-806-517
; Sequence 517, Application US/10102806
; Publication No. US20030054421A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies
; FILE REFERENCE: PA03P1C1
; CURRENT APPLICATION NUMBER: US/10/102,806
; PRIOR FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: 09/925,298
; PRIOR FILING DATE: 2001-08-10
; PRIOR APPLICATION NUMBER: PCT/US00/05881
; PRIOR FILING DATE: 2000-03-08
; PRIOR APPLICATION NUMBER: 60/124,270
; PRIOR FILING DATE: 1999-03-12
; NUMBER OF SEQ ID NOS: 846
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 517
; LENGTH: 250
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (118)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
; NAME/KEY: SITE
; LOCATION: (161)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
; NAME/KEY: SITE
; LOCATION: (204)
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids
US-10-102-806-517

Query Match 60.0%; Score 54; DB 9; Length 250;
Best Local Similarity 71.4%; Pred. No. 23;
Matches 10; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPR 14
||| ||| ||| |||
Db 202 RXHRPPAAPRRP 215

RESULT 6

US-09-030-619-96
; Sequence 96, Application US/09030619B

; Patent No. US20020035061A1
; GENERAL INFORMATION:
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erfile, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTIBIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030,619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 96
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Cationic Peptide Analogue
US-09-030-619-96

Query Match 58.9%; Score 53; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 2.6;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPPYLPFRPP 15
||| ||| ||| |||
Db 4 RPYIIPQPRP 14

RESULT 7

US-09-030-619-158
; Sequence 158, Application US/09030619B
; Patent No. US20020035061A1
; GENERAL INFORMATION:
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erfile, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTIBIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030,619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 158
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Apis mellifera
US-09-030-619-158

Query Match 58.9%; Score 53; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 2.6;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPPYLPFRPP 15
||| ||| ||| |||
Db 4 RPYIIPQPRP 14

RESULT 8

US-09-030-619-159
; Sequence 159, Application US/09030619B
; Patent No. US20020035061A1
; GENERAL INFORMATION:
; APPLICANT: Krieger, Timothy J.

```
; APPLICANT: Taylor, Robert
; APPLICANT: Efile, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INJECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTI-BIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030,619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 159
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Apis mellifera
US-09-030-619-159

Query Match      58.9%; Score 53; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 2.6;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYLPRPRPP 15
   |||:||||
Db 4 RPYIQPRPP 14

RESULT 9
US-09-030-619-160
; Sequence 160, Application US/09030619B
; Patent No. US20020035061A1
; GENERAL INFORMATION:
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Efile, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INJECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTI-BIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030,619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 160
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Apis mellifera
US-09-030-619-160

Query Match      57.8%; Score 52; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 3.3;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYLPRPRPP 15
   |||:||||
Db 4 RPYIQPRPP 14

RESULT 10
US-10-004-717-2
; Sequence 2, Application US/10004717
; Publication No. US20020192665A1
; GENERAL INFORMATION:
; APPLICANT: ZOGHBI, HUDA Y.
; APPLICANT: YANG, QI
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPEUTIC USE OF AN
; TITLE OF INVENTION: ATONAL ASSOCIATED SEQUENCE FOR DEAFNESS,
; TITLE OF INVENTION: OSTEOARTHRITIS, AND ABNORMAL CELL PROLIFERATION
; FILE REFERENCE: P01899US4
```

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; CURRENT APPLICATION NUMBER: US/10/004,717
; CURRENT FILING DATE: 2002-08-16
; PRIOR APPLICATION NUMBER: 09/585,645
; PRIOR FILING DATE: 2000-06-01
; PRIOR APPLICATION NUMBER: 60/176,993
; PRIOR FILING DATE: 2000-01-19
; PRIOR APPLICATION NUMBER: 60/137,060
; PRIOR FILING DATE: 1999-06-01
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 354
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-004-717-2

Query Match      57.8%; Score 52; DB 9; Length 354;
Best Local Similarity 57.1%; Pred. No. 52;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRPYPYLPYRPP 15
   |||:||||
Db 21 RQPQHLPQPPPP 34

RESULT 11
US-10-004-717-58
; Sequence 58, Application US/10004717
; Publication No. US20020192665A1
; GENERAL INFORMATION:
; APPLICANT: ZOGHBI, HUDA Y.
; APPLICANT: YANG, QI
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPEUTIC USE OF AN
; TITLE OF INVENTION: ATONAL ASSOCIATED SEQUENCE FOR DEAFNESS,
; FILE REFERENCE: P01899US4
; CURRENT APPLICATION NUMBER: US/10/004,717
; CURRENT FILING DATE: 2002-08-16
; PRIOR APPLICATION NUMBER: 09/585,645
; PRIOR FILING DATE: 2000-06-01
; PRIOR APPLICATION NUMBER: 60/176,993
; PRIOR FILING DATE: 2000-01-19
; PRIOR APPLICATION NUMBER: 60/137,060
; PRIOR FILING DATE: 1999-06-01
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 58
; LENGTH: 354
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-004-717-58

Query Match      57.8%; Score 52; DB 9; Length 354;
Best Local Similarity 57.1%; Pred. No. 52;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRPYPYLPYRPP 15
   |||:||||
Db 21 RQPQHLPQPPPP 34

RESULT 12
US-09-997-701-5
; Sequence 5, Application US/09997701
; Patent No. US20020107180A1
; GENERAL INFORMATION:
; APPLICANT: Yue, Henry
; APPLICANT: Corley, Neil C.
; APPLICANT: Guesler, Karl J.
; APPLICANT: Gorgone, Gina A.
; APPLICANT: Baughn, Mariah R.
; TITLE OF INVENTION: CELL SURFACE GLYCOPROTEINS
; FILE REFERENCE: PF-0631 US
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; CURRENT APPLICATION NUMBER: US/09/997,701
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/470,946
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-12-22
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: PERL Program
; SEQ ID NO 5
; LENGTH: 180
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: g2499136
US-09-997-701-5

Query Match 56.7%; Score 51; DB 10; Length 180;
Best Local Similarity 53.8%; Pred. No. 36;
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRPYPYLP-PRPP 15
:|:|:|:|:|:|
Db 47 KRPYPYQENP 59

RESULT 13

US-09-997-701-1
; Sequence 1, Application US/09997701
; Patent No. US20020107180A1
; GENERAL INFORMATION:
; APPLICANT: Yue, Henry
; APPLICANT: Corley, Neil C.
; APPLICANT: Guegler, Karl J.
; APPLICANT: Gorgone, Gina A.
; APPLICANT: Baughn, Mariah R.
; TITLE OF INVENTION: CELL SURFACE GLYCOPROTEINS
; FILE REFERENCE: PF-0631 US
; CURRENT APPLICATION NUMBER: US/09/997,701
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/470,946
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-12-22
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: PERL Program
; SEQ ID NO 1
; LENGTH: 195
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: 2297891
US-09-997-701-1

Query Match 56.7%; Score 51; DB 10; Length 195;
Best Local Similarity 53.8%; Pred. No. 39;
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRPYPYLP-PRPP 15
:|:|:|:|:|:|
Db 47 KRPYPYQENP 59

RESULT 14

US-09-747-835A-55
; Sequence 55, Application US/09747835A
; Patent No. US20020146692A1
; GENERAL INFORMATION:
; APPLICANT: Yamazaki, Victoria
; APPLICANT: Tang, Y. Tom
; APPLICANT: Liu, Chonghua
; APPLICANT: Zhou, Ping
; APPLICANT: Wang, Dunrui
; APPLICANT: Zhang, Jie
; APPLICANT: Ren, Feiyan
; APPLICANT: Asundi, Vinod
; APPLICANT: Drmanac, Radoje T
; TITLE OF INVENTION: METHODS AND MATERIALS RELATING TO G PROTEIN-COUPLED RECEPTOR-LIKE

; TITLE OF INVENTION: LIKE) POLYPEPTIDES AND POLYNUCLEOTIDES
; FILE REFERENCE: HYS-37CIP
; CURRENT APPLICATION NUMBER: US/09/747,835A
; CURRENT FILING DATE: 2002-03-08
; PRIOR APPLICATION NUMBER: US 09/729,739
; PRIOR FILING DATE: 2000-12-04
; PRIOR APPLICATION NUMBER: US 09/653,450
; PRIOR FILING DATE: 2000-08-31
; PRIOR APPLICATION NUMBER: US 09/620,312
; PRIOR FILING DATE: 2000-07-19
; PRIOR APPLICATION NUMBER: US 09/598,042
; PRIOR FILING DATE: 2000-06-20
; PRIOR APPLICATION NUMBER: US 09/552,317
; PRIOR FILING DATE: 2000-04-25
; PRIOR APPLICATION NUMBER: US 09/488,725
; PRIOR FILING DATE: 2000-01-21
; NUMBER OF SEQ ID NOS: 63
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 55
; LENGTH: 392
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-747-835A-55

Query Match 56.1%; Score 50.5; DB 10; Length 392;
Best Local Similarity 68.8%; Pred. No. 85;
Matches 11; Conservative 0; Mismatches 4; Indels 1; Gaps 1;

QY 1 RRRPRPYP-PRPP 15
||| ||| ||| |||
Db 368 RRRPNPRKPRGP 383

RESULT 15

US-10-243-035-2
; Sequence 2, Application US/10243035
; Publication No. US20030049697A1
; GENERAL INFORMATION:
; APPLICANT: LAZDUNSKI, MICHEL
; APPLICANT: LESAGE, FLORIAN
; APPLICANT: MAINGRET, FRANCOIS
; TITLE OF INVENTION: NEW FAMILY OF MECHANOSENSITIVE HUMAN POTASSIUM CHANNELS
; TITLE OF INVENTION: ACTIVATED BY POLYUNSATURATED FATTY ACIDS AND THEIR USE
; FILE REFERENCE: 1317-02
; CURRENT APPLICATION NUMBER: US/10/243,035
; CURRENT FILING DATE: 2002-09-13
; NUMBER OF SEQ ID NOS: 15
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 393
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-243-035-2

Query Match 56.1%; Score 50.5; DB 9; Length 393;
Best Local Similarity 68.8%; Pred. No. 85;
Matches 11; Conservative 0; Mismatches 4; Indels 1; Gaps 1;

QY 1 RRRPRPYP-PRPP 15
||| ||| ||| |||
Db 368 RRRPNPRKPRGP 383

Search completed: May 13, 2003, 10:42:32
Job time : 18 secs

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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:38:17 ; Search time 16 Seconds
(without alignments)
90.126 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPRPPYLPRPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 73:*

1: p1r1:*

2: p1r2:*

3: p1r3:*

4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	90	100.0	172	2 S68232	antimicrobial prot
2	66	73.3	59	2 A36589	bactenecin 7 - bov
3	59.5	66.1	82	2 A41051	spore coat protein
4	58	64.4	190	2 S68230	antimicrobial pept
5	56.5	62.8	168	2 S35330	apidaecin 14 precu
6	56.5	62.8	199	2 S14981	extensin class I (
7	55	61.1	437	2 A88942	protein R13D11.3 (
8	54.5	60.6	301	2 JQ1563	hybrid proline-ric
9	54	60.0	359	2 T13478	hypothetical prote
10	54	60.0	437	2 T32652	hypothetical prote
11	53	58.9	26	2 S06675	apidaecin 1b precu
12	53	58.9	144	2 S35331	apidaecin 22 precu
13	53	58.9	184	2 T29373	hypothetical prote
14	53	58.9	283	2 S35332	apidaecin 73 precu
15	53	58.9	428	2 E71415	probable coll wall
16	53	58.9	491	2 T07598	proline-rich prote
17	52	57.8	261	1 WMBEXE	infected cell prot
18	52	57.8	439	2 S19139	chitinase (EC 3.2.
19	52	57.8	467	2 S71169	protein kinase, 54
20	52	57.8	1066	2 G86292	hypothetical prote
21	51.5	57.2	1187	1 JC4155	protein-tyrosine-p
22	51.5	57.2	1189	1 JC2366	protein-tyrosine-p
23	51	56.7	180	2 S43791	PBDX protein - hum
24	50.5	56.1	1216	2 JW0105	synaptotaglin 2 alp
25	50	55.6	192	2 S76867	hypothetical prote
26	50	55.6	383	2 T06753	zinc finger protei
27	50	55.6	415	1 A34170	acrosin (EC 3.4.21
28	50	55.6	431	2 S29599	acrosin (EC 3.4.21
29	50	55.6	424	2 A54964	spliceosome-associ

30 50 55.6 449 2 S16748
31 50 55.6 547 2 S6828
32 50 55.6 1460 1 ED8E1F
33 50 55.6 3036 2 T18995
34 49.5 55.0 589 2 T29299
35 49 54.4 118 2 T19345
36 49 54.4 134 2 JC5572
37 49 54.4 161 2 F72593
38 49 54.4 210 2 T33700
39 49 54.4 218 2 T22261
40 49 54.4 296 2 A27319
41 49 54.4 296 2 S07361
42 49 54.4 352 2 F84799
43 49 54.4 369 2 S20500
44 49 54.4 380 2 T32944
45 49 54.4 413 2 H87604

proline-rich prote
unknown protein F1
immediate-early pr
hypothetical prote
hypothetical prote
hypothetical prote
proline-rich prote
hypothetical prote
hypothetical prote
gliadin - wheat
alpha/beta-gliadin
hypothetical prote
hydroxyproline-ric
hypothetical prote
hypothetical prote

ALIGNMENTS

RESULT 1

S68232

antimicrobial protein PR-39 precursor, cathelin-associated - pig

N:Alternate names: myeloid antibacterial protein PR-39

C:Species: Sus scrofa domestica (domestic pig)

C>Date: 15-Feb-1997 #sequence revision 13-Mar-1997 #text_change 20-Jun-2000

C:Accession: S68232, JN0899; I47138; S19563

R:Zhao, C.; Ganz, T.; Lehrer, R.I.

FEBS Lett. 376, 130-134, 1995

A:Title: Structures of genes for two cathelin-associated antimicrobial peptides: propheni

A:Reference number: S68232; MUID:96105365; PMID:7498526

A:Accession: S68232

A>Status: translation not shown

A:Molecule type: DNA

A:Residues: 1-172 <ZHA>

A:Cross-references: EMBL:X89201; NID:g1165150; PIDN:CAA61487.1; PID:g1165151

A:Experimental source: leukocytes

R:Storici, P.; Zanetti, M.

Biochem. Biophys. Res. Commun. 196, 1058-1065, 1993

A:Title: A cDNA derived from pig bone marrow cells predicts a sequence identical to the

A:Reference number: JN0899; MUID:94071853; PMID:8250863

A:Accession: JN0899

A:Molecule type: mRNA

A:Residues: 1-20, 'A', 22-172 <STO>

A:Cross-references: GB:L23825; NID:g435100; PIDN:AAA31109.1; PID:g435101

A:Experimental source: bone marrow cells

R:Gudmundsson, G.H.; Magnusson, K.P.; Chowdhary, B.P.; Johansson, M.; Andersson, L.; Bom

Proc. Natl. Acad. Sci. U.S.A. 92, 7085-7089, 1995

A:Title: Structure of the gene for porcine peptide antibiotic PR-39, a cathelin gene fami

A:Reference number: I47138; MUID:95350216; PMID:7624374

A:Accession: I47138

A>Status: preliminary; translated from GB/EMBL/DBBJ

A:Molecule type: DNA

A:Residues: 1-28, 'T', 30-89, 'QR', 92-116, 'NDP', 120-172 <GUD>

A:Cross-references: EMBL:X87336; NID:g829142; PIDN:CAA60682.1; PID:g1051298

R:Ageberth, B.; Lee, J.Y.; Bergman, T.; Carlquist, M.; Boman, H.G.; Mutt, V.; Joernvall,

Eur. J. Biochem. 202, 849-854, 1991

A:Title: Amino acid sequence of PR-39. Isolation from pig intestine of a new member of t

A:Reference number: S19563; MUID:92111534; PMID:1765098

A:Accession: S19563

A:Molecule type: protein

A:Residues: 131-169 <AGE>

A:Experimental source: intestine

C:Genetics:

A:Gene: PR39

A:Introns: 66/3; 102/3; 126/3

C:Superfamily: cathelin; cystatin homology

C:Keywords: amidated carboxyl end; antibacterial

F:1-29/Domain: signal sequence #status predicted <SIG>

F:22-129/Domain: cystatin homology <CYS>

F:30-130/Domain: propeptide #status predicted <PRO>

F:131-169/Product: antimicrobial protein PR-39 #status experimental <MAT>

P;169/Modified site: amidated carboxyl end (Pro) (amide in mature form from following gl

Query Match 100.0%; Score 90; DB 2; Length 172;
 Best Local Similarity 100.0%; Pred. No. 0.00035;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 15
 |||||
 DB 131 RRRPRPPVLPRLPRPP 145

RESULT 2
 A36589
 bacterescin 7 - bovine
 C;Species: Bos primigenius taurus (cattle)
 C;Date: 12-Apr-1991 #sequence_revision 12-Apr-1991 #text_change 09-May-1997
 C;Accession: A36589
 R;Frank, R.W.; Gennaro, R.; Schneider, K.; Przybylski, M.; Romeo, D.
 J. Biol. Chem. 265, 18871-18874, 1990
 A;Title: Amino acid sequences of two proline-rich bacterescins. Antimicrobial peptides of
 A;Reference number: A36589; MUID:91035404; PMID:2229048
 A;Accession: A36589
 A;Status: preliminary
 A;Molecule type: protein
 A;Residues: 1-59 <PRA>
 C;Superfamily: cathelin; cystatin homology

Query Match 73.3%; Score 66; DB 2; Length 59;
 Best Local Similarity 85.7%; Pred. No. 0.086;
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 14
 |||||
 DB 2:RIRPRPRLPRPP 15

RESULT 3
 A41051
 spore coat protein precursor - Bacillus subtilis
 C;Species: Bacillus subtilis
 C;Date: 03-Apr-1992 #sequence_revision 06-Jan-1995 #text_change 11-Jan-2002
 C;Accession: S04835; A41051; F69606
 R;Aranson, A.I.; Song, H.Y.; Bourne, N.
 Mol. Microbiol. 3, 437-444, 1989
 A;Title: Gene structure and precursor processing of a novel Bacillus subtilis spore coat
 A;Reference number: S04835; MUID:89313296; PMID:2546006
 A;Accession: S04835
 A;Molecule type: DNA
 A;Residues: 'MNVHTPLNSIRNMVGIKKAREVFL', 2-82 <AR2>
 A;Cross-references: EMBL:X13740; NID:G39864; PIDN:CAA32004.1; PID:G39865
 A;Experimental source: strain JH642
 A;Note: Part of this sequence, including the amino end of the mature protein, was confir
 R;Bourne, N.; FitzJames, P.C.; Aranson, A.I.
 J. Bacteriol. 173, 6618-6625, 1991
 A;Title: Structural and germination defects of Bacillus subtilis spores with altered con
 A;Reference number: A41051; MUID:92011439; PMID:1917883
 A;Accession: A41051
 A;Molecule type: protein
 A;Residues: 'XX', 3-11 <BOU>
 A;Experimental source: strain JH642
 A;Note: the material sequenced was the larger of two isolated precursor forms; the amino
 A;Note: both the location of the transcription start site and peptide sequencing of the
 R;Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Berter
 C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Chd
 A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Brington, J.; Fabret, C.; Ferrari, E.
 Nature 390, 249-256, 1997
 A;Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Galizzi, A.; Gallen
 iech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holsappel, S.; Hosono, S.; Hullo, M.F.
 Koetter, P.; Konigstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois
 A;Authors: Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Muegel
 Y, M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portetelle
 Rieger, M.; Rivolta, C.; Rocha, E.; Roche, M.; Rose, M.; Sadaie, Y.; Sato, T.; Scanlon,
 A;Authors: Schleich, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seron

akeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Tosato, V.; Uchiyama,
 T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, K
 A;Authors: Yoshikawa, H.F.; Zumstein, E.; Yoshikawa, H.; Danchin, A.
 A;Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.
 A;Reference number: A69580; MUID:98044033; PMID:9384377
 A;Accession: F69606
 A;Status: preliminary; nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 'MNVHTPLNSIRNMVGIKKAREVFL', 2-82 <KUN>
 A;Cross-references: GB:Z99110; GB:AL009126; NID:G2633472; PIDN:CAB13066.1; PID:G2633563
 A;Experimental source: strain 168
 C;Comment: This structural protein is expressed during stage V of sporulation.
 C;Genetics:
 A;Gene: cotT
 A;Start codon: TTG
 C;Keywords: sporulation
 F;1-19/Domain: propeptide #status experimental <PRO>
 F;20-82/Product: spore coat protein #status experimental <MAT>

Query Match 66.1%; Score 59.5; DB 2; Length 82;
 Best Local Similarity 84.6%; Pred. No. 0.69;
 Matches 11; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 4 PRPP-YLPRLPRPP 15
 |||||
 DB 49 PRPPYYPRPP 61

RESULT 4
 S68230
 antimicrobial peptide precursor - sheep
 N;Alternate names: Bac7.5 peptide homolog
 C;Species: Ovis orientalis aries, Ovis ammon aries (domestic sheep)
 C;Date: 15-Feb-1997 #sequence_revision 13-Mar-1997 #text_change 21-Jan-2000
 C;Accession: S68230
 R;Bagella, L.; Scocchi, M.; Zanetti, M.
 FEBS Lett. 376, 225-228, 1995
 A;Title: cDNA sequences of three sheep myeloid cathelicidins.
 A;Reference number: S68228; MUID:96105386; PMID:7498547
 A;Accession: S68230
 A;Status: preliminary
 A;Molecule type: mRNA
 A;Residues: 1-190 <BAG>
 A;Cross-references: EMBL:I46852; NID:g1161244; PIDN:AA85468.1; PID:g1161245
 C;Superfamily: cathelin; cystatin homology
 F;1-23/Domain: signal sequence #status predicted <SIG>
 F;22-129/Domain: signal sequence #status predicted <SIG>
 F;29-130/Domain: cystatin homology <CYS>
 F;130-190/Product: propeptide #status predicted <PRO>
 F;130-190/Product: antimicrobial peptide #status predicted <MAT>

Query Match 64.4%; Score 58; DB 2; Length 190;
 Best Local Similarity 78.6%; Pred. No. 2.4;
 Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 14
 |||||
 DB 132 RLRPRPRLPRPP 145

RESULT 5
 S35330
 apidaecin 14 precursor - honeybee
 N;Contains: apidaecin II
 C;Species: Apis mellifera (honeybee)
 C;Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 21-Jul-2000
 C;Accession: S35330; S06676
 R;Casteels-Josson, K.; Capaci, T.; Casteels, P.; Tempst, P.
 EMBO J. 12, 1569-1578, 1993
 A;Title: Apidaecin multipetide precursor structure: a putative mechanism for amplificati
 A;Reference number: S35330; MUID:93223697; PMID:8467807
 A;Accession: S35330
 A;Molecule type: mRNA
 A;Residues: 1-168 <CAS>

A;Residues: 1-427 <CHI>
 A;Cross-references: EMBL:AF039043; PIDN:AAB94196.1; GSPDB:GNO0028; CESP:F39C12.3
 A;Experimental source: strain Bristol N2; clone F39C12
 C;Genetics:
 A;Gene: CESP:F39C12.3
 A;Map position: X
 A;Introns: 42/3; 104/3; 133/3; 164/3; 213/3; 276/3; 336/3

Query Match 60.0%; Score 54; DB 2; Length 427;

Best Local Similarity 69.2%; Pred. No. 16;
 Matches 9; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRPYPYLRPRPP 15

|||||.|||

Db 338 RRPDPDIPPLPP 350

RESULT 11

S06675

apidaecin 1b precursor - honeybee

C;Species: Apis mellifera (honeybee)

C;Date: 07-Sep-1990 #sequence_revision 07-Sep-1990 #text_change 16-Dec-1998

C;Accession: S06675

R;Casteels, P.; Ampe, C.; Jacobs, F.; Vaeck, M.; Tempst, P.

EMBO J. 8, 2387-2391, 1989

A;Title: Apidaecins: antibacterial peptides from honeybees.

A;Reference number: S05383; MUID:90005446; PMID:2676519

A;Accession: S06675

A;Molecule type: protein

A;Residues: 1-26 <CAS>

F;1-8/Domain: propeptide #status experimental <PRO>

F;9-26/Product: apidaecin 1b #status experimental <MAT>

Query Match 58.9%; Score 53; DB 2; Length 26;

Best Local Similarity 72.7%; Pred. No. 1.3;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLRPRPP 15

|||||.|||

Db 12 RPVIYQPRPP 22

RESULT 12

S35331

apidaecin 22 precursor - honeybee

C;Species: Apis mellifera (honeybee)

C;Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 21-Jul-2000

C;Accession: S35331

R;Casteels-Josson, K.; Capaci, T.; Casteels, P.; Tempst, P.

EMBO J. 12, 1569-1578, 1993

A;Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplification

A;Reference number: S35330; MUID:93223697; PMID:8467807

A;Accession: S35331

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-144 <CAS>

A;Cross-references: EMBL:X72576; NID:9297064; PIDN:CAA51168.1; PID:9297065

C;Superfamily: procyclic acidic repetitive protein

Query Match 58.9%; Score 53; DB 2; Length 144;

Best Local Similarity 72.7%; Pred. No. 7.1;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLRPRPP 15

|||||.|||

Db 102 RPVIYQPRPP 112

RESULT 13

T29373

hypothetical protein ZC404.1 - Caenorhabditis elegans

C;Species: Caenorhabditis elegans

C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jun-2000

C;Accession: T29373
 R;Bentley, D.; Le, T.T.
 submitted to the EMBL Data Library, April 1996
 A;Description: The sequence of C. elegans cosmid ZC404.

A;Reference number: Z20614

A;Accession: T29373

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-184 <BEN>

A;Cross-references: EMBL:U55363; PIDN:AAA97967.1; GSPDB:GNO0023; CESP:ZC404.1

A;Experimental source: strain Bristol N2; clone ZC404

C;Genetics:

A;Gene: CESP:ZC404.1

A;Map position: 5

A;Introns: 15/2; 50/2; 75/2; 138/2

C;Superfamily: Caenorhabditis elegans hypothetical protein ZC404.1

Query Match 58.9%; Score 53; DB 2; Length 184;

Best Local Similarity 90.0%; Pred. No. 9;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 RRPYPYLRPP 12

|||||.|||

Db 26 RPRKPYLRPP 35

RESULT 14

S35332

apidaecin 73 precursor - honeybee (fragment)

N;Contains: apidaecin 1a

C;Species: Apis mellifera (honeybee)

C;Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 03-Nov-2000

C;Accession: S35332; S05383

R;Casteels-Josson, K.; Capaci, T.; Casteels, P.; Tempst, P.

EMBO J. 12, 1569-1578, 1993

A;Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplification

A;Reference number: S35330; MUID:93223697; PMID:8467807

A;Accession: S35332

A;Molecule type: mRNA

A;Residues: 1-283 <CAS>

A;Cross-references: EMBL:X72577; NID:9297066; PIDN:CAA51169.1; PID:94539289

A;Accession: S05383

A;Molecule type: protein

A;Residues: 258-283 <CA3>

C;Superfamily: proline-rich protein

F;266-283/Product: apidaecin 1a #status experimental <MAT>

Query Match 58.9%; Score 53; DB 2; Length 283;

Best Local Similarity 72.7%; Pred. No. 14;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLRPRPP 15

|||||.|||

Db 241 RPVIYQPRPP 251

RESULT 15

E71415

probable coll wall protein - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

A;Variety: Columbia

C;Date: 03-Aug-1998 #sequence_revision 03-Aug-1998 #text_change 05-Dec-1998

C;Accession: E71415

R;Bevan, M.; Bancroft, I.; Bent, E.; Love, K.; Goodman, H.; Dean, C.; Bergkamp, R.; Dirks

P.; Wedler, H.; Wedler, E.; Wambutt, R.; Weitzenecker, T.; Pohl, T.M.; Terry, N.; Giele

avanagh, T.; Hempel, S.; Kotter, P.; Entian, K.D.; Rieger, M.; Schaeffer, M.; Funk, B.

Nature 391, 485-488, 1998

A;Authors: Mueller-Auer, S.; Silvey, M.; James, R.; Montfort, A.; Pons, A.; Puigdomenech,

erhoft, A.; Moores, T.; Jones, J.D.G.; Eneva, T.; Palme, K.; Benes, V.; Rechman, S.; Ansc

C.; Chalwatzis, N.

A;Title: Analysis of 1.9 Mb of contiguous sequence from chromosome 4 of Arabidopsis thali

A;Reference number: A71400; MUID:98121113; PMID:9461215

A;Accession: E71415

A;Status: preliminary; nucleic acid sequence not shown; translation not shown

A;Molecule type: DNA

A;Residues: 1-428 <BEV>

A;Cross-references: GB:Z97338; NID:G2244870; PID:G327461; PID:G224487*

C;Genetics:

A;Map position: 4COP9-4G3845

Query Match 58.9%; Score 53; DB 2; Length 428;
Best Local Similarity 61.5%; Pred. No. 21;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 3 RRPYPYLPYRPP 15

Db 67 KPPPPYIPCPPP 79

Search completed: May 13, 2003, 10:41:48
Job time : 18 secs

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GenCore version 5.1.4 p5 4578
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:14:02 ; Search time 11 seconds
(without alignments)
56.559 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPRPPYLPRPP 15

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_40.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	90	100.0	172	1 PR39_PIG	P80054 sus scrofa
2	66	73.3	190	1 BCT7_BOVIN	P19661 bos taurus
3	59.5	66.1	107	1 COTT_BACSU	P11863 bacillus su
4	58	64.4	190	1 BCT7_SHEEP	P50415 ovis aries
5	56.5	62.8	168	1 AP14_APIME	Q06601 apis mellif
6	55.5	61.7	151	1 RNB_HSV2H	P89479 herpes simp
7	53	58.9	144	1 AP22_APIME	P35581 apis mellif
8	53	58.9	283	1 PRLP_BOVIN	Q06602 apis mellif
9	53	58.9	381	1 PRLP_BOVIN	Q9gkx8 bos taurus
10	52	57.8	261	1 RLI_HSV2H	P28283 herpes simp
11	52	57.8	354	1 ATHI_HUMAN	Q92858 homo sapien
12	52	57.8	467	1 AFCI_ARATH	P51566 arabidopsis
13	52	57.8	841	1 RELA_STRAT	Q85709 streptomyce
14	51.5	57.2	1187	1 PTNE_HUMAN	Q15678 homo sapien
15	51.5	57.2	1189	1 PTNE_MOUSE	Q62130 mus musculu
16	51	56.7	15	1 MK1_PALPR	P80408 palomela pr
17	51	56.7	180	1 XG_HUMAN	P55808 homo sapien
18	50.5	56.1	393	1 C1W4_HUMAN	Q9ny98 homo sapien
19	50	55.6	17	1 APID_BOMPA	P81464 bombus pasc
20	50	55.6	415	1 ACRO_PIG	P08001 sus scrofa
21	50	55.6	424	1 S3B4_HUMAN	Q15427 homo sapien
22	50	55.6	449	1 APG_BRANA	P40603 brassica na
23	50	55.6	678	1 ABPE_RIPCL	Q27905 riptortus c
24	49	54.4	134	1 PRL5_HUMAN	Q99954 homo sapien
25	49	54.4	296	1 GDA6_WHEAT	P04726 triticum ae
26	49	54.4	352	1 RRS1_ARATH	Q9sh88 arabidopsis
27	49	54.4	2911	1 FBN2_HUMAN	P35556 homo sapien
28	48.5	53.9	2142	1 BAT2_HUMAN	P48634 homo sapien
29	48	53.3	280	1 TNF6_CERTO	Q9bdn1 cercocebus
30	48	53.3	280	1 TNF6_MACMU	Q9myl6 macaca mla
31	48	53.3	281	1 TNF6_HUMAN	P48023 homo sapien
32	48	53.3	282	1 TNF6_PIG	Q9bea8 sus scrofa
33	48	53.3	402	1 VGLD_PRVRI	P07645 pseudorabie

RESULT 1

ID	PR39_PIG	STANDARD;	PRT;	172 AA.
AC	P80054; Q9TR84;			
DT	01-MAR-1992 (Rel. 21, Created)			
DT	01-OCT-1996 (Rel. 34, Last sequence update)			
DT	16-OCT-2001 (Rel. 40, Last annotation update)			
DE	Antibacterial protein PR-39 precursor.			
GN	PR39.			
OS	Sus scrofa (Pig).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.			
OX	NCBI_TaxID=9823;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=95350216; PubMed=7624374;			
RA	Gudmundsson G.H., Magnusson K.P., Chowdhary B.P., Johansson M.,			
RA	Andersson L., Boman H.G.;			
RT	"Structure of the gene for porcine peptide antibiotic PR-39, a			
RT	cathelin gene family member: comparative mapping of the locus for the			
RT	human peptide antibiotic FALL-39."			
RL	Proc. Natl. Acad. Sci. U.S.A. 92:7085-7089(1995).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Bone marrow;			
RX	MEDLINE=94071853; PubMed=8250863;			
RA	Storici P., Zanetti M.;			
RT	"A cDNA derived from pig bone marrow cells predicts a sequence			
RT	identical to the intestinal antibacterial peptide PR-39."			
RL	Biochem. Biophys. Res. Commun. 196:1058-1065(1993).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Liver;			
RX	MEDLINE=96105365; PubMed=7498526;			
RA	Zhao C., Ganz T., Lehrer R.I.;			
RT	"Structures of genes for two cathelin-associated antimicrobial			
RT	peptides: prophenin-2 and PR-39."			
RL	FEBS Lett. 376:130-134(1995).			
RN	[4]			
RP	SEQUENCE OF 131-169.			
RC	TISSUE=Intestine;			
RX	MEDLINE=92111534; PubMed=1765098;			
RA	Agarberth B., Lee J.-Y., Bergman T., Carlquist M., Boman H.G.,			
RA	Mutt V., Joernvall H.;			
RT	"Amino acid sequence of PR-39. Isolation from pig intestine of a new			
RT	member of the family of proline-arginine-rich antibacterial			
RT	peptides."			
RL	Eur. J. Biochem. 202:849-854(1991).			
RN	[5]			
RP	SEQUENCE OF 131-164, AND FUNCTION.			
RC	TISSUE=Neutrophils;			
RX	MEDLINE=95088504; PubMed=7996056;			
RA	Shi J., Ross C.R., Chengappa M.M., Blecha F.;			
RT	"Identification of a proline-arginine-rich antibacterial peptide from			
RT	neutrophils that is analogous to PR-39, an antibacterial peptide from			
RT	the small intestine."			

ALIGNMENTS

34	48	53.3	520	1	C84A_ARATH	Q42600 arabidopsis
35	48	53.3	759	1	TOP3_CAEEL	O61660 caenorhabdi
36	47.5	52.8	283	1	EXTN_SORBI	P24152 sorghum bic
37	47.5	52.8	372	1	DBFA_HUMAN	P16989 homo sapien
38	47.5	52.8	1443	1	SYO2_HUMAN	O15056 homo sapien
39	47	52.2	176	1	BCT5_BOVIN	P19660 bos taurus
40	47	52.2	261	1	PRP2_MOUSE	P05142 mus musculu
41	47	52.2	296	1	PRP3_MOUSE	P05143 mus musculu
42	47	52.2	507	1	MEFA_HUMAN	Q02078 homo sapien
43	47	52.2	753	1	SK30_HUMAN	O94993 homo sapien
44	47	52.2	846	1	IR31_HCMVA	P09715 human cytom
45	47	52.2	1040	1	BO12_YEAST	P39969 saccharomyc

```

RL J. Leukoc. Biol. 56:807-811(1994).
CC -1- FUNCTION: EXERTS A POTENT ANTIMICROBIAL ACTIVITY AGAINST BOTH
CC E. COLI AND B. MEGATERIUM.
CC -1- TISSUE SPECIFICITY: SMALL INTESTINE AND BONE MARROW.
CC -1- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.
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CC -----
DR EMBL; X87236; CAA60682.1; -.
DR EMBL; L23825; AAA31109.1; -.
DR EMBL; X89201; CAA61487.1; -.
DR PIR; S19563; S19563.
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidin; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
KW Antibiotic; Amidation; Signal.
FT SIGNAL 1 29
FT PROPEP 30 130
FT CHAIN 131 169
FT MOD_RES 30 30
FT DISULFID 85 96
FT DISULFID 107 124
FT MOD_RES 169 169
FT CONFLICT 21 21
FT CONFLICT 29 29
FT CONFLICT 90 91
FT CONFLICT 117 119
FT CONFLICT 157 157
FT SEQUENCE 172 AA; 19476 MW; 994B792798C0E133 CRC64;
SQ
Query Match 100.0%; Score 90; DB 1; Length 172;
Best Local Similarity 100.0%; Pred. No. 0.00022;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 RRRPRPPYLP RPRPP 15
Db 131 RRRPRPPYLP RPRPP 145
RESULT 2
ID BCT7_BOVIN STANDARD; PRT; 190 AA.
AC P19661;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Bactenecin 7 precursor (BAC7) (PR-59).
GN BAC7.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=95010707; PubMed=7925973;
RA Scocchi M., Romeo D., Zanetti M.;
RT "Molecular cloning of Bact7, a proline- and arginine-rich
RT antimicrobial peptide from bovine neutrophils.";
RL FEBS Lett. 352:197-200(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;

```

```

RA Scocchi M., Wang S., Zanetti M.;
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE OF 131-189.
RC TISSUE=Neutrophils;
RX MEDLINE=91035404; PubMed=2229048;
RA Frank R.W., Gennaro R., Schneider K., Przybylski M., Romeo D.;
RT "Amino acid sequences of two proline-rich bactericins. Antimicrobial
RT peptides of bovine neutrophils.";
RL J. Biol. Chem. 265:18871-18874(1990).
RN [4]
RP CHARACTERIZATION.
RX MEDLINE=96300243; PubMed=8706679;
RA Storici P., Tossi A., Lenarcic B., Romeo D.;
RT "Purification and structural characterization of bovine
RT cathelicidins, precursors of antimicrobial peptides.";
RL Eur. J. Biochem. 238:769-776(1996).
CC -1- FUNCTION: EXERTS, IN VITRO, A POTENT ANTIMICROBIAL ACTIVITY.
CC PROBABLY DUE TO AN IMPAIRMENT OF THE FUNCTION OF THE RESPIRATORY
CC CHAIN AND OF ENERGY-DEPENDENT ACTIVITIES IN THE INNER MEMBRANE
CC OF SUSCEPTIBLE MICROORGANISMS.
CC -1- TISSUE SPECIFICITY: LARGE GRANULES OF NEUTROPHILS.
CC -1- PTM: ELASTASE IS RESPONSIBLE FOR ITS MATURATION.
CC -1- MASS SPECTROMETRY: MW=18395; MW_ERR=1; METHOD=Electrospray;
CC RANGE=30-190.
CC -1- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.
CC -----
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CC -----
DR EMBL; L42977; AAA87359.1; -.
DR EMBL; Y0471; CAA70616.1; -.
DR PIR; A36589; A36589.
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidin; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
KW Antibiotic; Repeat; Signal.
FT SIGNAL 1 29
FT PROPEP 30 130
FT CHAIN 131 190
FT MOD_RES 189 190
FT DISULFID 85 96
FT DISULFID 107 124
FT SEQUENCE 190 AA; 21567 MW; 8CD07D7AA30A731C CRC64;
SQ
Query Match 73.3%; Score 66; DB 1; Length 190;
Best Local Similarity 85.7%; Pred. No. 0.15;
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 RRRPRPPYLP RPRPP 14
Db 132 RRRPRPPYLP RPRPP 145
RESULT 3
COTT_BACSU STANDARD; PRT; 107 AA.
ID COTT_BACSU
AC P11863;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Spore coat protein T precursor.
GN COTT.
OS Bacillus subtilis.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.

```



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OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=168 / JH642;
RX MEDLINE=89313296; PubMed=2546006;
RA Aronson A.I., Song H.Y., Bourne N.;
RT "Gene structure and precursor processing of a novel Bacillus subtilis
  spore coat protein.";
RL Mol. Microbiol. 3:437-444 (1989).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=168;
RX MEDLINE=98044033; PubMed=9384377;
RA Kunst F., Ogasawara N., Moszer I., Albertini A.M., Alloni G.,
  Azevedo V., Bertero M.G., Bessieres P., Bolotin A., Borchert S.,
  Borriss R., Boursier L., Brans A., Braun M., Brignell S.C., Bron S.,
  Brouillet S., Bruschini C.V., Caldwell B., Capuano V., Carter N.M.,
  Choi S.K., Codani J.J., Conneron I.F., Cummings N.D., Daniel N.R.A.,
  Denizot F., Devine K.M., Dusterhoft A., Ehrlich S.D., Emmerson P.T.,
  Entia K.D., Errington J., Fabret C., Ferrari E., Foulger D.,
  Fritz C., Fujita M., Fujita Y., Funa S., Galizzi A., Galleron N.,
  Chim S.Y., Glaser P., Goffeau A., Gollightly E.J., Grandi G.,
  Guisepi G., Gay B.J., Haga K., Haeck J., Harwood C.R., Henaut A.,
  Hilbert H., Holsappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,
  Joris B., Karamata D., Kasahara Y., Klaer-Blanchard M., Klein C.,
  Kobayashi Y., Koetter P., Koningstein G., Krogh S., Kumano M.,
  Kurita K., Lapidus A., Lardinois S., Lauber J., Lazarevic V.,
  Lee S.M., Levine A., Liu H., Masuda S., Mael C., Medigue C.,
  Medina N., Mellado R.P., Mizuno M., Moestl D., Nakai S., Noback M.,
  Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,
  Parro V., Pohl T.M., Portetelle D., Porwollik S., Prescott A.M.,
  Pressac E., Pujic P., Purnelle B., Rapoport G., Rey M., Reynolds S.,
  Rieger M., Rivolta C., Rocha E., Roche B., Rose M., Sadaie Y.,
  Sato T., Scanlan E., Schleich S., Schroeter P., Scoffone F.,
  Sekiguchi J., Sekowska A., Seror S.J., Serror P., Shin B.S., Soldo B.,
  Sorokin A., Tacconi E., Takagi T., Takahashi H., Takemaru K.,
  Takeuchi M., Tamakoshi A., Tanaka T., Terpstra P., Tognoni A.,
  Tosato V., Uchiyama S., Vandenbol M., Vannier F., Vassarotti A.,
  Viari A., Wambitt R., Wedler E., Wedler H., Weitzensegger T.,
  Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
  Yoshida K., Yoshikawa H.F., Zumbstein E., Yoshikawa H., Danchin A.;
RT "The complete genome sequence of the Gram-positive bacterium Bacillus
  subtilis.";
RL Nature 390:249-256 (1997).
CC -1- FUNCTION: POSSIBLY PROTECTION OF SPORE AND PROBABLY PLAYS
  SOME ROLE IN GERMINATION.
CC -1- SUBCELLULAR LOCATION: OUTER SURFACE OF ENDOSPORE.
CC
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DR EMBL; X13740; CAA32004.1; -
DR EMBL; Z99110; CAB13066.1; -
DR FIR; S04835; S04835.
DR Subtilisin; BG10495; cotT.
KW Sporulation; Signal; Complete proteome.
FT SIGNAL 1 44
FT CHAIN 45 107 SPORE COAT PROTEIN T.
SQ SEQUENCE 107 AA; 12992 MW; AD1P66F0C4CE29A3 CRC64;

Query Match 66.1%; Score 59.5; DB 1; Length 107;
Best Local Similarity 84.6%; Pred. No. 0.47;
Matches 11; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 4 PRPP-YLPRPRPP 15
DB 74 PRPPYYPRPRPP 86
|||||

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```

RESULT 4
BCT7_SHEEP STANDARD; PRT; 190 AA.
ID AC P50415;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Bactenein 7 precursor (BAC7).
GN BAC7.5.
OS Ovis aries (Sheep).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Caprinae; Ovis.
OC NCBI_TaxID=9940;
RN [1]_TaxID=9940;
RP SEQUENCE FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=96105386; PubMed=7498547;
RA Bagella L., Scocchi M., Zanetti M.;
RT "cDNA sequences of three sheep myeloid cathelicidins.";
RL FEBS Lett. 376:225-228 (1995).
CC -1- FUNCTION: EXERTS, IN VITRO, A POTENT ANTIMICROBIAL ACTIVITY.
  PROBABLY DUE TO AN IMPAIRMENT OF THE FUNCTION OF THE RESPIRATORY
  CHAIN AND OF ENERGY-DEPENDENT ACTIVITIES IN THE INNER MEMBRANE
  OF SUSCEPTIBLE MICROORGANISMS (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.
CC
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  or send an email to license@isb-sib.ch).
CC
DR EMBL; L46852; AAA85468.1; -
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidin; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
KW Antibiotic; Repeat; Signal.
FT SIGNAL 1 29 POTENTIAL.
FT PROPEP 30 130 BY SIMILARITY.
FT CHAIN 131 190 PYRROLIDONE CARBOXYLIC ACID
FT MOD_RES 30 30
FT DISULFID 85 96 BY SIMILARITY.
FT DISULFID 107 124 BY SIMILARITY.
SQ SEQUENCE 190 AA; 21829 MW; E4AAFB1600E98371 CRC64;

Query Match 64.4%; Score 58; DB 1; Length 190;
Best Local Similarity 78.6%; Pred. No. 1.2;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RRRPRPPYLRPRPP 14
DB 132 RLRRPRRLPRPRPP 145
|||||

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RESULT 5
API4_APIME STANDARD; PRT; 168 AA.
ID AC Q06601; P11525; P11526; P11527;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE Apidaecin precursor, type 14.
GN APID14.
OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
OC Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;

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```
OC ACuleata; Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;
RN SEQUENCE FROM N.A.
RP MEDLINE=93223697; PubMed=8467807;
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RX "Apidaecin multipeptide precursor structure: a putative mechanism for
RT amplification of the insect antibacterial response.";
RL EMBO J. 12:1569-1578(1993).
RN [2]
RP SEQUENCE OF APIDAECINS IA/IB/II.
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMBO J. 8:2387-2391(1989).
CC -1- FUNCTION: APIDAECINS HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY
CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
CC PROPAGATION.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL: X72575; CAA51167.1; -.
DR PIR: S05383; S05383.
DR PIR: S06675; S06675.
DR PIR: S06676; S06676.
DR PIR: S35330; S35330.
DR InterPro: IPR004828; Apidaecin.
DR Pfam: PF00807; Apidaecin; 5.
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
KW Cleavage on pair of basic residues; Repeat.
FT SIGNAL 1 19
FT PROPEP 35 42
FT PEPTIDE 43 60
FT PROPEP 63 70
FT PEPTIDE 71 88
FT PROPEP 91 98
FT PEPTIDE 99 116
FT PROPEP 119 124
FT PEPTIDE 125 142
FT PROPEP 145 150
FT PEPTIDE 151 168
SQ SEQUENCE 168 AA; 19380 MW; 594B931254C04A37 CRC64;
Query Match 62.8%; Score 56.5; DB 1; Length 168;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 11; Conservative 2; Mismatches 2; Indels 7; Gaps 1;
QY 1 RRRP-----RPPYLPRLPRPP 15
DB 117 RREPEAPGNRRPVYIQPRPP 138
RESULT 6
RNB_HSV2H
ID_RNB_HSV2H STANDARD; PRT; 151 AA.
AC P89479;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Potential RNA-binding protein.
GN US11.
OS Herpes simplex virus (type 2 / strain HG52).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Simplexvirus.
OX NCBI_TaxID=10315;
RN [1]
RP SEQUENCE FROM N.A.
RA Dolan A.;
RL Submitted (FEB-1997) to the EMBL/GenBank/DBSJ databases.
CC -1- FUNCTION: BINDS DNA AND RNA (BY SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar (By similarity).
CC
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CC
CC EMBL: Z86099; CAB06719.1; -.
KW DNA-binding; RNA-binding; Repeat; Nuclear protein.
FT DOMAIN 90 146
FT REPEAT 90 95
FT REPEAT 96 101
FT REPEAT 102 104
FT REPEAT 105 110
FT REPEAT 111 116
FT REPEAT 117 122
FT REPEAT 123 128
FT REPEAT 129 130
FT REPEAT 131 134
FT REPEAT 135 140
FT REPEAT 141 146
SQ SEQUENCE 151 AA; 16297 MW; FAB751F23C3DB6AE CRC64;
Query Match 61.7%; Score 55.5; DB 1; Length 151;
Best Local Similarity 73.3%; Pred. No. 1.9;
Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;
QY 2 RRPRLPRLPR-PRPP 15
DB 127 RPPRLPRLPRPRPP 141
RESULT 7
AP22_APIME
ID AP22_APIME STANDARD; PRT; 144 AA.
AC P35581; P11525; P11526;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE Apidaecin precursor, type 22.
OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
OC Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;
OC Aculeata; Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=93223697; PubMed=8467807;
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RX "Apidaecin multipeptide precursor structure: a putative mechanism for
RT amplification of the insect antibacterial response.";
RL EMBO J. 12:1569-1578(1993).
RN [2]
RP SEQUENCE (APIDAECINS IA/IB).
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMBO J. 8:2387-2391(1989).
CC -1- FUNCTION: APIDAECINS HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY
CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
CC PROPAGATION.
CC
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 CC -----

DR EMBL; X72576; CAA51168.1; -.
 DR PIR; S05383; S05383.
 DR PIR; S06675; S06675.
 DR PIR; S35331; S35331.
 DR InterPro; IPR004828; Apidaecin.
 DR Pfam; PF00807; Apidaecin; 4.
 KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
 KW Cleavage on pair of basic residues; Repeat.
 FT SIGNAL 1 19 POTENTIAL.
 FT PROPEP 35 42
 FT PEPTIDE 43 60 APIDAEACIN IB.
 FT PROPEP 63 70
 FT PEPTIDE 71 88 APIDAEACIN IB.
 FT PROPEP 91 98
 FT PEPTIDE 99 116 APIDAEACIN IB.
 FT PROPEP 119 126
 FT PEPTIDE 127 144 APIDAEACIN IA.
 SQ SEQUENCE 144 AA; 16539 MW; 6FAIAD74CB77108D CRC64;

Query Match 58.9%; Score 53; DB 1; Length 144;
 Best Local Similarity 72.7%; Pred. No. 3.6;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 5 RPYLPRLPRPP 15

DB 102 RPVIYIQQRP 112
 |||:|:|:|

RESULT 8

AP73_APIME STANDARD; PRT; 283 AA.
 AC Q06602; P11525; P11526;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 01-JUN-1994 (Rel. 29, Last annotation update)
 DE Apidaecin precursor, type 73 (Fragment).
 GN APID73.
 OS Apis mellifera (Honeybee).
 OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
 OC Insecta; Pserygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;
 OC Aculeata; Apoidea; Apidae; Apis.
 OX NCBI_TaxID=7460;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93223697; PubMed=8467807;
 RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
 RT "Apidaecin multipetide precursor structure: a putative mechanism for
 RT amplification of the insect antibacterial response.";
 RL EMBO J. 12:1569-1578(1993).
 RN [2]
 RP SEQUENCE OF APIDAEACIN IA/IB.
 RC TISSUE=Hemolymph;
 RX MEDLINE=90005446; PubMed=2676519;
 RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
 RT "Apidaecins: antibacterial peptides from honeybees.";
 RL EMBO J. 8:2387-2391(1989).
 CC -!- FUNCTION: APIDAEACIN HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY
 CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
 CC PROPAGATION.
 CC -----

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 CC -----

DR EMBL; X72577; CAA51169.1; -.
 DR PIR; S05383; S05383.
 DR PIR; S06675; S06675.
 DR PIR; S35332; S35332.
 DR InterPro; IPR004828; Apidaecin.
 DR Pfam; PF00807; Apidaecin; 9.
 KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
 KW Cleavage on pair of basic residues; Repeat.
 FT NON_TER 1 18
 FT SIGNAL <1 18 POTENTIAL.
 FT PROPEP 34 41 APIDAEACIN IB.
 FT PEPTIDE 42 59
 FT PROPEP 62 69
 FT PEPTIDE 70 87 APIDAEACIN IB.
 FT PROPEP 90 97
 FT PEPTIDE 98 115 APIDAEACIN.
 FT PROPEP 118 125
 FT PEPTIDE 126 143 APIDAEACIN IB.
 FT PROPEP 146 153
 FT PEPTIDE 154 171 APIDAEACIN.
 FT PROPEP 174 182
 FT PEPTIDE 183 199 APIDAEACIN IB.
 FT PROPEP 202 209
 FT PEPTIDE 210 227 APIDAEACIN IB.
 FT PROPEP 230 237
 FT PEPTIDE 238 255 APIDAEACIN IB.
 FT PROPEP 258 265
 FT PEPTIDE 266 283 APIDAEACIN IA.
 SQ SEQUENCE 283 AA; 32695 MW; 4EA5FDECD5E142B CRC64;

Query Match 58.9%; Score 53; DB 1; Length 283;
 Best Local Similarity 72.7%; Pred. No. 6.9;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 5 RPYLPRLPRPP 15

DB 241 RPVIYIQQRP 251
 |||:|:|:|

RESULT 9

PRLP_BOVIN STANDARD; PRT; 381 AA.
 ID PRLP_BOVIN
 AC Q9GKN8;
 DT 15-JUN-2002 (Rel. 41, Created)
 DT 15-JUN-2002 (Rel. 41, Last sequence update)
 DT 15-JUN-2002 (Rel. 41, Last annotation update)
 DE Proargin precursor (Proline-arginine-rich end leucine-rich repeat
 DE of protein).
 GN PRELP.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Articular cartilage;
 RX MEDLINE=20576219; PubMed=11007795;
 RA Bengtsson E., Asberg A., Heinegaard D., Sommarin Y., Spillmann D.;
 RT "The amino-terminal part of PRELP binds to heparin and heparan
 RT sulfate.";
 RL J. Biol. Chem. 275:40695-40702(2000).
 RN [2]
 RP FUNCTION.
 RX PubMed=11847210;
 RA Bengtsson E., Moergelin M., Sasaki T., Timpl R., Heinegaard D.,
 RA Asberg A.;
 RT "The leucine-rich repeat protein PRELP binds perlecan and collagens
 RT and may function as a basement membrane anchor.";
 RL J. Biol. Chem. 277:15061-15068(2002).
 CC -!- FUNCTION: May anchor basement membranes to the underlying
 CC connective tissue.
 CC -!- SUBUNIT: Binds the basement membrane heparan sulfate proteoglycan

CC perlecan and triple helical collagens type I and type II.
CC -1- SUBCELLULAR LOCATION: Extracellular matrix.
CC -1- DOMAIN: The basic amino-terminal Arg/Pro-rich binds heparin and
CC heparan sulfate. Binds collagen type I and type II through its
CC leucine-rich repeat domain.
CC -1- SIMILARITY: BELONGS TO THE SMALL LEUCINE-RICH PROTEOGLYCANS
CC (SLRPs) FAMILY. CLASS II SUBFAMILY.
CC -1- SIMILARITY: CONTAINS 12 LEUCINE-RICH REPEATS (LRR).
CC
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CC
CC EMBL: AF163568; AAG23723.1; -
CC InterPro: IPR001611; LRR.
CC InterPro: IPR003372; LRR Nterm.
CC InterPro: IPR003592; LRR Out.
CC InterPro: IPR003591; LRR Typ.
CC Pfam: PF00560; LRR; 10.
CC Pfam: PF01462; LRRNT; 1.
CC PRINTS: PR00019; LEURICHRPT.
CC SMART: SM00370; LRR; 7.
CC SMART: SM00013; LRRNT; 1.
CC SMART: SM00369; LRR Typ; 7.
CC Glycoprotein; Extracellular matrix; Repeat; Leucine-rich repeat;
CC Signal.
KW SIGNAL.
FT CHAIN 1 21 POTENTIAL.
FT PROLARGIN 22 381
FT CYS-RICH 72 88
FT DOMAIN 94 113
FT REPEAT 114 137 LRR-T 1.
FT REPEAT 138 161 LRR-T 2.
FT REPEAT 162 182 LRR-S 2.
FT REPEAT 183 206 LRR-T 3.
FT REPEAT 207 232 LRR-T 4.
FT REPEAT 233 253 LRR-S 3.
FT REPEAT 254 277 LRR-T 5.
FT REPEAT 278 302 LRR-T 6.
FT REPEAT 303 322 LRR-S 4.
FT REPEAT 323 361 LRR-T 7.
FT REPEAT 362 381 LRR-T 8.
FT DOMAIN 196 201 POLY-LEU.
FT DISULFID 331 372 BY SIMILARITY.
FT CARBOHYD 123 123 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 288 288 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 319 319 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 381 AA; 43682 MW; 23DA99C01B772A0 CRC64;

Query Match 58.9%; Score 53; DB 1; Length 381;
Best Local Similarity 76.9%; Pred. No. 9.2;
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 RRRPRPPVLRPRRP 14
DB 25 RRRPRPPVLRPRRP 37
|||||
RESULT 10
RL1_HSV2H STANDARD; PRT; 261 AA.
AC P28283;
DT 01-DEC-1992 (Rel. 24, Created)
DT 01-DEC-1992 (Rel. 24, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Neurovirulence factor (ICP34.5).
GN RL1.
OS Herpes simplex virus (type 2 / strain HG52).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;

OC Alphaherpesvirinae; Simplexvirus.
OX NCBI_TaxID=10315;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=92113549; PubMed=1662697;
RA McGeech D.J., Cunningham C., McIntyre G., Dolan A.;
RT "Comparative sequence analysis of the long repeat regions and
RT adjoining parts of the long unique regions in the genomes of herpes
RT simplex viruses types 1 and 2";
RL J. Gen. Virol. 72:3057-3075(1991).
RN [2]
RP SEQUENCE FROM N.A.
RA Dolan A.;
RL Submitted (FEB-1997) to the EMBL/GenBank/DBJ databases.
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CC
CC EMBL: D10471; BAA23428.1; -
CC EMBL: Z86099; CAB06759.1; -
CC EMBL: Z86099; CAB06706.1; -
CC PIR: JQ1502; WMBEXE.
KW Repeat.
FT DOMAIN 3 12 2 X 5 AA TANDEM REPEATS OF R-R-R-G-P.
FT REPEAT 3 7
FT REPEAT 8 12 2 X 8 AA TANDEM REPEATS OF P-R-P-G-A-P-A-
FT DOMAIN 16 31 V.
FT REPEAT 16 23
FT REPEAT 24 31
FT SEQUENCE 261 AA; 27908 MW; 4BBD13AF3D906D71 CRC64;

Query Match 57.8%; Score 52; DB 1; Length 261;
Best Local Similarity 64.7%; Pred. No. 8.3;
Matches 11; Conservative 1; Mismatches 3; Indels 2; Gaps 1;

QY 1 RRRPRP--PYLPRPRRP 15
DB 13 RRRPRPGAPVPRPGAP 29
||||| :|||
RESULT 11
ATH1_HUMAN STANDARD; PRT; 354 AA.
ID ATH1_HUMAN
AC Q92858;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE Atonal protein homolog 1 (Helix-loop-helix protein hATH-1).
GN ATH1 OR ATH1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97026280; PubMed=8872459;
RA Ben-Arie N., McCall A.E., Berkman S., Eichele G., Bellen H.J.,
RA Zoghbi H.Y.;
RT "Evolutionary conservation of sequence and expression of the bHLH
RT protein Atonal suggests a conserved role in neurogenesis.";
RL Hum. Mol. Genet. 5:1207-1216(1996).
CC -1- FUNCTION: ACTIVATES E BOX-DEPENDENT TRANSCRIPTION IN COLLABORATION
CC WITH E47, BUT THE ACTIVITY IS COMPLETELY ANTAGONIZED BY THE
CC NEGATIVE REGULATOR OF NEUROGENESIS HES-1. MAY PLAY A ROLE IN THE
CC DIFFERENTIATION OF SUBSETS OF NEURAL CELLS BY ACTIVATING E BOX-
CC DEPENDENT TRANSCRIPTION (BY SIMILARITY).
CC -1- SUBUNIT: EFFICIENT DNA BINDING REQUIRES DIMERIZATION WITH ANOTHER

```
CC BHLH PROTEIN.
CC -!- SUBCELLULAR LOCATION: Nuclear (Probable).
CC -!- SIMILARITY: BELONGS TO THE BASIC HELIX-LOOP-HELIX (BHLH) FAMILY OF
CC TRANSCRIPTION FACTORS. "ATONAL" SUBFAMILY.
CC -----
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CC -----
CC EMBL; U61148; AAB41305.1; -.
CC TRANSFAC; T04544; -.
CC Genew; HGNC:797; ATOH1.
CC MIM; 601461; -.
CC InterPro; IPR001092; HLH_basic.
CC Pfam; PF00010; HLH; 1.
CC SMART; SM00353; HLH; 1.
CC PROSITE; PS00038; HLH_1; FALSE_NEG.
CC Transcription regulation; Activator; DNA-binding; Nuclear protein.
KW DOMAIN 29 38 POLY-PRO.
FT DNA_BIND 160 171 BASIC DOMAIN.
FT DOMAIN 172 212 HELIX-LOOP-HELIX MOTIF (BY SIMILARITY).
FT DOMAIN 224 228 POLY-PRO.
SQ SEQUENCE 354 AA; 39160 MW; AB12F1E917A00A8D CRC64;

Query Match 57.8%; Score 52; DB 1; Length 354;
Best Local Similarity 57.1%; Pred. NO. 11;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRRPPYLPRLPPRP 15
DB 21 *RQPHLPQPPPP 34

RESULT 12
AFCL_ARATH STANDARD; PRT; 467 AA.
AC P51566;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Protein kinase AFCL (EC 2.7.1.1.-).
GN AFCL.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopses.
OX NCBI_TaxID=3702;
RN [1]
SEQUENCE FROM N.A.
RC STRAIN=cv. Landsberg erecta;
RX MEDLINE=95083650; PubMed=7991592;
RA Bender J., Fink G.R.;
RT dependent processes in yeast.;
RL Proc. Natl. Acad. Sci. U.S.A. 91:12105-12109(1994).
CC -!- FUNCTION: ACTIVATOR OF YEAST TRANSCRIPTION FACTOR, STE12.
CC -!- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
CC LAMMER SUBFAMILY.
CC -----
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CC -----
CC EMBL; U16176; AAA57117.1; -.

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DR HSP; P24941; 1AQ1.
DR InterPro; IPR000719; Euk_pkinase.
DR InterPro; IPR002290; Ser_thr_pkinase.
DR Pfam; PF00669; pkinase; 1.
DR ProDom; PD000001; Euk_pkinase; 1.
DR SMART; SM00220; S_TK; 1.
DR PROSITE; PS00107; PROTEIN KINASE_ATP; FALSE_NEG.
DR PROSITE; PS00108; PROTEIN KINASE_ST; 1.
DR PROSITE; PS50011; PROTEIN KINASE_DOM; 1.
KW Transferase; Serine/threonine-protein kinase; ATP-binding.
FT DOMAIN 115 443 PROTEIN KINASE.
FT NP_BIND 121 129 ATP (BY SIMILARITY).
FT BINDING 144 144 ATP (BY SIMILARITY).
FT ACT_SITE 240 240 BY SIMILARITY.
SQ SEQUENCE 467 AA; 54216 MW; 54D739A82F490E12 CRC64;

Query Match 57.8%; Score 52; DB 1; Length 467;
Best Local Similarity 52.4%; Pred. NO. 15;
Matches 11; Conservative 1; Mismatches 3; Indels 6; Gaps 1;

QY 1 RRRPR-----PPYLPRLPPRP 15
DB 35 RRRPRLTWDRAAPPLPPPPPP 55

RESULT 13
RELA_STRAT STANDARD; PRT; 841 AA.
ID RELA_STRAT
AC O85709;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE GTP pyrophosphokinase (EC 2.7.6.5) (ATP:GTP 3'-pyrophosphotransferase)
DE (PPGPP synthetase I) (PPGPP synthetase).
GN RELA.
OS Streptomyces antibioticus.
OC Bacteria; Actinobacteria; Actinobacteria (class); Actinobacteridae;
OC Actinomycetales; Streptomycetales; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=1890;
RN [1]
SEQUENCE FROM N.A.
RC STRAIN=IMRU 3720;
RX MEDLINE=99296594; PubMed=10368159;
RA Hoyt S., Jones G.H.;
RT "relA is required for actinomycin production in Streptomyces
RT antibioticus.";
RL J. Bacteriol. 181:3824-3829(1999).
CC -!- FUNCTION: In eubacteria PPGPP (guanosine 3'-diphosphate 5'-
CC diphosphate) is a mediator of the stringent response that
CC coordinates a variety of cellular activities in response to
CC changes in nutritional abundance. This enzyme catalyzes the
CC formation of PPGPP which is then hydrolysed to form PPGPP (By
CC similarity). Is required for actinomycin production.
CC -!- CATALYTIC ACTIVITY: ATP + GTP = AMP + guanosine 3'-diphosphate 5'-
CC triphosphate.
CC -!- PATHWAY: FIRST STEP IN THE METABOLISM OF PPGPP.
CC -!- SIMILARITY: BELONGS TO THE RELA / SPOT FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF072829; AAC26021.1; -.
DR InterPro; IPR002912; ACT.
DR InterPro; IPR002819; HD.
DR InterPro; IPR003607; ME_Ppase_Hdc.
DR InterPro; IPR004811; Spot_rela.
DR InterPro; IPR004095; TGS_dom.
DR Pfam; PF01842; ACT; 1.

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DR Pfam; PF01966; HD; 1.
DR Pfam; PF02824; TGS; 1.
DR SMART; SM00471; HDC; 1.
DR TIGRFAMs; TIGR00691; spot_rela; 1.
SQ SEQUENCE 841 AA; 93671 MW; 632A037BA4BF4C94 CRC64;

Query Match 57.8%; Score 52; DB 1; Length 841;
Best Local Similarity 60.0%; Pred. No. 26;
Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 1 RRRPPYLPRLPRPP 15
   |||||
Db 50 RPKPAPPRPPPP 64

RESULT 14
PTNE HUMAN STANDARD; PRT; 1187 AA.
AC Q15678;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Protein tyrosine phosphatase, non-receptor type 14 (EC 3.1.3.48)
DE (Protein-tyrosine phosphatase pez).
GN PTPN14 OR PEZ.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Breast carcinoma;
RX MEDLINE=95251727; PubMed=7733990;
RA Smith A.L., Mitchell P.J., Shipley J., Gusterson B.A., Rogers M.V.,
RA Crompton M.R.;
RT "Pez": a novel human cDNA encoding protein tyrosine phosphatase- and
RT ezrin-like domains."
RL Biochem. Biophys. Res. Commun. 209:959-965(1995).
CC -1- CATALYTIC ACTIVITY: Protein tyrosine phosphatase + H(2)O = protein
CC tyrosine + phosphate.
CC -1- TISSUE SPECIFICITY: EXPRESSED IN A VARIETY OF HUMAN TISSUES
CC INCLUDING KIDNEY, SKELETAL MUSCLE, LUNG AND PLACENTA.
CC -1- SIMILARITY: CONTAINS 1 BAND 4.1-LIKE DOMAIN.
CC -1- SIMILARITY: BELONGS TO THE NON-RECEPTOR CLASS OF THE PROTEIN-
CC TYROSINE PHOSPHATASE FAMILY.
-----
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-----
DR EMBL; X82676; CAA57993.1; -
DR HSSP; P29350; LGWZ.
DR Genew; HGNC:9647; PTPN14.
DR MIM; 603155; -
DR InterPro; IPR000299; Band 4.1.
DR InterPro; IPR000387; TYR_Pp.
DR InterPro; IPR000242; TYR_PP.
DR Pfam; PF00102; Y_phosphatase; 1.
DR Pfam; PF00373; Band 41; 1.
DR PRINTS; PR00935; BAND41.
DR PRINTS; PR00700; PRTYPHPTASE.
DR SMART; SM00194; PTPC; 1.
DR SMART; SM00295; B41; 1.
DR PROSITE; PS00660; BAND_41_1; 1.
DR PROSITE; PS00661; BAND_41_2; 1.
DR PROSITE; PS00657; BAND_41_3; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS00055; TYR_PHOSPHATASE_PTP; 1.

DR PROSITE; PS00056; TYR_PHOSPHATASE_2; 1.
DR Structural protein; Cytoskeleton; Hydrolase.
DR DOMAIN 25 239 BAND 4.1-LIKE.
DR DOMAIN 933 1187 PROTEIN-TYROSINE PHOSPHATASE.
DR FT ACT SITE 1121 1121 BY SIMILARITY.
DR FT DOMAIN 566 573 POLY-PRO.
DR FT DOMAIN 709 716 POLY-GLU.
SQ SEQUENCE 1187 AA; 135239 MW; 015760B75E3574E3 CRC64;

Query Match 57.2%; Score 51.5; DB 1; Length 1187;
Best Local Similarity 83.3%; Pred. No. 42;
Matches 10; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 3 RRRPPYLPRLPRPP 14
   |||||
Db 565 RPPPPY-PRPRP 575

RESULT 15
PTNE MOUSE STANDARD; PRT; 1189 AA.
AC Q62130;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Protein tyrosine phosphatase, non-receptor type 14 (EC 3.1.3.48)
DE (Protein-tyrosine phosphatase PTP36).
GN PTPN14.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CB-17-SCID; TISSUE=Thymus;
RX MEDLINE=94354945; PubMed=8074693;
RA Sawada M., Ogata M., Fujino Y., Hamakata T.;
RT "cDNA cloning of a novel protein tyrosine phosphatase with homology
RT to cytoskeletal protein 4.1 and its expression in T-lineage cells."
RL Biochem. Biophys. Res. Commun. 203:479-484(1994).
CC -1- FUNCTION: MAY BE INVOLVED IN THE REGULATION OF T CELL DEVELOPMENT.
CC -1- CATALYTIC ACTIVITY: Protein tyrosine phosphatase + H(2)O = protein
CC tyrosine + phosphate.
CC -1- TISSUE SPECIFICITY: THYMUS; IN CELLS OF BOTH HEMATOPOIETIC AND
CC NON-HEMATOPOIETIC ORIGINS.
CC -1- SIMILARITY: CONTAINS 1 BAND 4.1-LIKE DOMAIN.
CC -1- SIMILARITY: BELONGS TO THE NON-RECEPTOR CLASS OF THE PROTEIN-
CC TYROSINE PHOSPHATASE FAMILY.
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DR EMBL; D31842; BAA06628.1; -
DR HSSP; Q06124; ZSHP.
DR MGD; MGI:102467; Ptpn14.
DR InterPro; IPR000299; Band 4.1.
DR InterPro; IPR000387; TYR_Pp.
DR InterPro; IPR000242; TYR_PP.
DR Pfam; PF00102; Y_phosphatase; 1.
DR Pfam; PF00373; Band 41; 1.
DR PRINTS; PR00935; BAND41.
DR PRINTS; PR00700; PRTYPHPTASE.
DR SMART; SM00295; B41; 1.
DR SMART; SM00194; PTPC; 1.
DR PROSITE; PS00660; BAND_41_1; 1.
DR PROSITE; PS00661; BAND_41_2; 1.
DR PROSITE; PS00657; BAND_41_3; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS00055; TYR_PHOSPHATASE_PTP; 1.

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DR PROSITE; PS50055; TYR PHOSPHATASE PTP; 1.
DR PROSITE; PS50056; TYR PHOSPHATASE 2; 1.
KW Structural protein; Cytoskeleton; Hydrolase.
FT DOMAIN 75 239 BAND 4.1-LIKE.
FT DOMAIN 935 1189 PROTEIN-TYROSINE PHOSPHATASE.
FT ACT SITE 1123 1123 BY SIMILARITY.
FT DOMAIN 566 573 POLY-PRO.
FT DOMAIN 635 639 POLY-GLY.
FT DOMAIN 712 718 POLY-GLU.
SQ SEQUENCE 1189 AA; 135030 MW; 2B85BE5F9C723303 CRC64;

Query Match 57.2%; Score 51.5; DB 1; Length 1189;
Best Local Similarity 83.3%; Pred. No. 42;
Matches 10; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

OY 3 RPPPPYLP RPP 14
DB 565 RPPPPY-P RPP 575

Search completed: May 13, 2003, 10:40:51
Job time : 13 secs

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GenCore version 5.1.4.p5.4578
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:37:02 ; Search time 29 seconds
(without alignments)
106.576 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPPPPYLRPRPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 671580 seqs, 206047115 residues

Total number of hits satisfying chosen parameters: 671580

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_21.*

- 1: sp archaea.*
- 2: sp bacteria.*
- 3: sp fungi.*
- 4: sp human.*
- 5: sp invertebrate.*
- 6: sp mammal.*
- 7: sp mhc.*
- 8: sp organelle.*
- 9: sp phase.*
- 10: sp plant.*
- 11: sp rodent.*
- 12: sp virus.*
- 13: sp vertebrate.*
- 14: sp unclassified.*
- 15: sp rvirus.*
- 16: sp bacteriaph.*
- 17: sp archesp.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	61	67.8	336	12 Q68405	Q68405 human cytom
2	58	64.4	156	10 Q8RV32	Q8RV32 oryza sativ
3	58	64.4	164	6 P79361	P79361 ovis aries
4	58	64.4	190	6 Q9X3Q9	Q9X3Q9 capra hircu
5	58	64.4	224	6 O19031	O19031 ovis aries
6	57.5	63.9	183	10 Q94J98	Q94J98 oryza sativ
7	57	63.3	200	16 Q9RK54	Q9RK54 streptomyce
8	57	63.3	361	2 Q9XCG4	Q9XCG4 mycobacteri
9	56	62.2	212	2 O08306	O08306 nocardioid
10	55	61.1	212	12 O41980	O41980 murid herpe
11	55	61.1	464	12 Q91TM2	Q91TM2 tupaia herp
12	54.5	60.6	301	10 Q41848	Q41848 zea mays (m
13	54.5	60.6	2635	12 P88955	P88955 kaposi's sa
14	54.5	60.6	2635	12 O40942	O40942 kaposi's sa
15	54	60.0	347	11 Q8R353	Q8R353 mus musculu
16	54	60.0	359	5 Q9XZT0	Q9XZT0 drosophila

17	54	60.0	427	5	O44582	O44582 caenorhabdi
18	54	60.0	451	5	Q95X63	Q95X63 caenorhabdi
19	54	60.0	955	4	Q9Y2W1	Q9Y2W1 homo sapien
20	53.5	59.4	225	11	Q99JA6	Q99JA6 mus musculu
21	53	58.9	168	10	Q9SM77	Q9SM77 oryza sativ
22	53	58.9	184	5	Q23291	Q23291 caenorhabdi
23	53	58.9	185	10	Q94JF6	Q94JF6 oryza sativ
24	53	58.9	199	5	Q8WSY8	Q8WSY8 apis mellif
25	53	58.9	333	10	Q9X123	Q9X123 oryza sativ
26	53	58.9	428	10	Q23370	Q23370 arabidopsis
27	53	58.9	491	10	O82066	O82066 solanum tub
28	53	58.9	520	10	Q9LV14	Q9LV14 arabidopsis
29	52.5	58.3	602	12	Q66852	Q66852 fowl adenov
30	52	57.8	148	16	Q8U5T2	Q8U5T2 agrobacteri
31	52	57.8	155	4	Q96E55	Q96E55 homo sapien
32	52	57.8	439	10	O42421	O42421 beta vulgar
33	52	57.8	450	10	Q94CE1	Q94CE1 arabidopsis
34	52	57.8	467	10	Q39184	Q39184 arabidopsis
35	52	57.8	790	5	Q8T458	Q8T458 drosophila
36	52	57.8	1006	10	Q9LMQ1	Q9LMQ1 arabidopsis
37	52	57.8	1091	5	Q9W1Z6	Q9W1Z6 drosophila
38	51.5	57.2	145	12	Q8V718	Q8V718 simian herp
39	51.5	57.2	238	10	Q8W097	Q8W097 oryza sativ
40	51	56.7	94	5	Q917F1	Q917F1 drosophila
41	51	56.7	145	10	Q8SAY8	Q8SAY8 oryza sativ
42	51	56.7	255	10	Q8RYX5	Q8RYX5 oryza sativ
43	51	56.7	409	5	Q9U0Z7	Q9U0Z7 leishmania
44	51	56.7	417	5	Q9V4V1	Q9V4V1 drosophila
45	51	56.7	470	10	Q8S503	Q8S503 oryza sativ

ALIGNMENTS

RESULT 1

Q68405 PRELIMINARY; PRT; 336 AA.
AC Q68405;
DT 01-NOV-1996 (T-EMBLrel. 01, Created)
DT 01-NOV-1996 (T-EMBLrel. 01, Last sequence update)
DT 01-NOV-1998 (T-EMBLrel. 08, Last annotation update)
DE Orf UL151.
OS Human cytomegalovirus.
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Betaherpesvirinae; Cytomegalovirus.
OX NCBI_TaxID=10359;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=TOLEDO;
RX MEDLINE=96099416; PubMed=8523595;
RA Cha T.A., Tom E., Kemble G.W., Duke G.M., Mocarski E.S., Spaete R.R.;
RT "Human cytomegalovirus clinical isolates carry at least 19 genes not
found in laboratory strains.";
RL J. Virol. 70:78-83(1996).
DR EMBL; U33331; AAA8582.1; --
SQ SEQUENCE 336 AA; 35116 MW; 9F865E5019F69D0C CRC64;

Query Match 67.8%; Score 61; DB 12; Length 336;
Best Local Similarity 78.6%; Pred. No. 0.94;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 RRRPPPPYLRPRPP 15

Db 279 RRRPPIQLQRRPP 292

RESULT 2

ID Q8RV32 PRELIMINARY; PRT; 156 AA.
AC Q8RV32;
DT 01-JUN-2002 (T-EMBLrel. 21, Created)
DT 01-JUN-2002 (T-EMBLrel. 21, Last sequence update)
DT 01-JUN-2002 (T-EMBLrel. 21, Last annotation update)

```

DE OSJNB0032K15.1 protein (OJ1159_D09.32 protein).
GN OSJNB0032K15.1 OR OJ1159_D09.32.
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzeae; Oryza.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. NIPPONBARE;
RA Sasaki T., Matsumoto T., Yamamoto K.;
RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC
clone:OSJNB0032K15.";
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. NIPPONBARE;
RA Sasaki T., Matsumoto T., Yamamoto K.;
RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC
clone:OJ1159_D09.";
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP003710; BAB86560.1; -.
DR EMBL; AP003792; BAB89214.1; -.
SQ SEQUENCE 156 AA; 17659 MW; 4152112C3DB493CF CRC64;

Query Match 64.4%; Score 58; DB 10; Length 156;
Best Local Similarity 73.3%; Pred. No. 1.2;
Matches 11; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRP 15
DB 78 RRRPRPPYLP RPRP 92

RESULT 3
ID P79361 PRELIMINARY; PRT; 164 AA.
AC P79361;
DT 01-MAY-1997 (TrEMBLrel. 03, Created)
DT 01-MAY-1997 (TrEMBLrel. 03, Last sequence update)
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
DE 7.5 kDa bactinecin (Fragment).
GN BAC7.5.
OS Ovis aries (Sheep).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Caprinae; Ovis.
OX NCBI_TaxID=9940;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=LIVER;
RX MEDLINE=96140581; PubMed=8549789;
RA Mahoney M.M., Lee A.Y., Brezinski-Caliguri D.J., Huttner K.M.;
RT "Molecular analysis of the sheep cathelin family reveals a novel
antimicrobial peptide.";
RL FEBS Lett. 377:519-522(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=LIVER;
RA Huttner K.M., Mahoney M.M.;
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; U60598; AAB49713.1; -.
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidins; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
FT NON TER 164
SQ SEQUENCE 164 AA; 16642 MW; E3BFC871F6AE8B9A CRC64;

Query Match 64.4%; Score 58; DB 6; Length 164;
Best Local Similarity 78.6%; Pred. No. 1.2;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY 1 RRRPRPPYLP RPRP 14
DB 132 RLRPRRPLRPRP 145

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RESULT 4

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O9XSQ9 PRELIMINARY; PRT; 190 AA.
AC O9XSQ9;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE BAC7.5 protein.
GN BAC7.5.
OS Capra hircus (Goat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Caprinae; Capra.
OX NCBI_TaxID=9925;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=BONE MARROW;
RA Zhao C., Nguyen T., Brogden K., Lehrer R.;
RT "cDNA cloning of goat cathelin related peptides.";
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ243125; CAB45523.1; -.
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidins; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
FT CHAIN 131
SQ SEQUENCE 190 AA; 21835 MW; D13305EF16875F4F CRC64;

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Query Match 64.4%; Score 58; DB 6; Length 190;
Best Local Similarity 78.6%; Pred. No. 1.4;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY 1 RRRPRPPYLP RPRP 14
DB 132 RLRPRRPLRPRP 145

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RESULT 5

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O19031 PRELIMINARY; PRT; 224 AA.
AC O19031;
DT 01-JAN-1998 (TrEMBLrel. 05, Created)
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)
DE BACTINECIN 11 precursor.
GN BAC11.
OS Ovis aries (Sheep).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Caprinae; Ovis.
OX NCBI_TaxID=9940;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98121317; PubMed=9461419;
RA Huttner K.M., Lambeth M.R., Burkin H.R., Broad T.E.;
RT "Localization and genomic organization of sheep antimicrobial peptides
genes.";
RL Gene 206:85-91(1998).
CC -!- FUNCTION: ANTIMICROBIAL PEPTIDE (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.
DR EMBL; U77049; AAB62000.1; -.
DR EMBL; U77046; AAB62000.1; JOINED.
DR EMBL; U77047; AAB62000.1; JOINED.
DR EMBL; U77048; AAB62000.1; JOINED.
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.

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DR ProDom; PD001838; Cathelicidin; 1.
 DR PROSITE; PS00946; CATHELICIDINS 1; 1.
 DR PROSITE; PS00947; CATHELICIDINS 2; 1.
 KW Signal; Antibiotic.
 FT SIGNAL 1 29 POTENTIAL.
 FT PROPEP 30 130 POTENTIAL.
 FT CHAIN 131 224 BACTINECIN 11.
 FT MOD_RES 30 30 PYROLIDONE CARBOXYLIC ACID (BY
 FT SIMILARITY).
 FT DISULFID 85 96 BY SIMILARITY.
 FT DISULFID 107 124 BY SIMILARITY.
 SQ SEQUENCE 224 AA; 25669 MW; 6AEAB1256AC76FC CRC64;
 Query Match 64.4%; Score 58; DB 6; Length 224;
 Best Local Similarity 78.6%; Pred. No. 1.6;
 Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLRPRP 14
 |||||
 DB 132 RLRRPRRLPRPP 145
 RESULT 6
 Q94J98 PRELIMINARY; PRT; 183 AA.
 AC Q94J98;
 DT 01-DEC-2001 (TrEMBLrel. 19, Created)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DE P0047B08.14 protein (OJ1159.D09.5 protein).
 GN P0047B08.14 OR OJ1159.D09.5.
 OS Oryza sativa (Rice), and
 OS Oryza sativa (japonica cultivar-group).
 OS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzeae; Oryza.
 OX NCBI_TaxID=4530, 39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. NIPPONBARE;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, PAC
 RT clone:P0047B08."
 RL Submitted (DEC-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. NIPPONBARE;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC
 RT clone:OJ1159.D09."
 RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AP003053; BAB55690.1;
 DR EMBL; AP003792; BAB89188.1;
 SQ SEQUENCE 183 AA; 20155 MW; F1CF82AD89CEB36 CRC64;
 Query Match 63.9%; Score 57.5; DB 10; Length 183;
 Best Local Similarity 73.3%; Pred. No. 1.6;
 Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;
 QY 1 RRRPRPPYLRPRPP 15
 |||||
 DB 129 RSRPR-PYAPRPQP 142
 RESULT 7
 Q9RK54 PRELIMINARY; PRT; 200 AA.
 AC Q9RK54;
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DE P0047B08.14 protein (OJ1159.D09.5 protein).
 GN P0047B08.14 OR OJ1159.D09.5.
 OS Oryza sativa (Rice), and
 OS Oryza sativa (japonica cultivar-group).
 OS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzeae; Oryza.
 OX NCBI_TaxID=4530, 39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. NIPPONBARE;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, PAC
 RT clone:P0047B08."
 RL Submitted (DEC-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CV. NIPPONBARE;
 RA Sasaki T., Matsumoto T., Yamamoto K.;
 RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC
 RT clone:OJ1159.D09."
 RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AP003053; BAB55690.1;
 DR EMBL; AP003792; BAB89188.1;
 SQ SEQUENCE 183 AA; 20155 MW; F1CF82AD89CEB36 CRC64;
 Query Match 63.9%; Score 57.5; DB 10; Length 183;
 Best Local Similarity 73.3%; Pred. No. 1.6;
 Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;
 QY 1 RRRPRPPYLRPRPP 15
 |||||
 DB 129 RSRPR-PYAPRPQP 142

OS Streptomyces coelicolor.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.
 OX NCBI_TaxID=1902;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A3(2) / M145;
 RA Bentley S.D., Chater K.F., Cerdano-Tarraga A.-M., Challis G.L.,
 RA Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,
 RA Harper D., Bateman A., Brown S., Chandra G., Hornsby T., Collins M.,
 RA Cronin A., Fraser A., Goble L., Hidaigo J., Hensby T., O'Neil S.,
 RA Huang C.-H., Kieser T., Larke L., Murphy L., Oliver K., O'Neil S.,
 RA Rabinowitz E., Rajandream M.A., Rutherford K., Rutter S.,
 RA Seeger K., Saunders D., Sharp S., Squares R., Squares S., Taylor K.,
 RA Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,
 RA Hopwood D.A.;
 RT "Complete genome sequence of the model actinomycete Streptomyces
 RT coelicolor A3(2)."
 RL Nature 417:141-147(2002).
 DR EMBL; AL117669; CAB56128.1;
 KW Hypothetical protein.
 SQ SEQUENCE 200 AA; 22076 MW; 0DCBBEC5585803B5 CRC64;
 Query Match 63.3%; Score 57; DB 16; Length 200;
 Best Local Similarity 76.9%; Pred. No. 2;
 Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLRPRP 13
 |||||
 DB 118 RRHPEPPALPRP 130
 RESULT 8
 Q9XCG4 PRELIMINARY; PRT; 361 AA.
 ID Q9XCG4;
 AC Q9XCG4;
 DT 01-NOV-1999 (TrEMBLrel. 12, Created)
 DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE Hypothetical 40.2 kDa protein.
 OS Mycobacterium avium.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1764;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=2151;
 RA Eckstein T.M., Lambert M.L., Brennan P.J., Belisle J.T., Inamine J.M.;
 RT "Identification of a gene cluster involved in glycopeptidolipid
 RT biosynthesis and of a gene cluster encoding daunorubicin resistance in
 RT two strains of Mycobacterium avium serovar 2."
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AF143772; AAD44199.1;
 KW Hypothetical protein.
 SQ SEQUENCE 361 AA; 40208 MW; AD01DBE825C1C9EA CRC64;
 Query Match 63.3%; Score 57; DB 2; Length 361;
 Best Local Similarity 71.4%; Pred. No. 3.4;
 Matches 10; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
 QY 1 RRRPRPPYLRPRP 14
 |||||
 DB 32 RRRPRPPAPHPPP 45
 RESULT 9
 O08306 PRELIMINARY; PRT; 212 AA.
 ID O08306;
 AC O08306;
 DT 01-JUL-1997 (TrEMBLrel. 04, Created)
 DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE Hypothetical 22.7 kDa protein.

OS Nocardioideae simplex (Arthrobacter simplex).
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Propionibacterineae; Nocardioideaceae; Pimelobacter.
OX NCBI_TaxID=2045;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IFO12069;
RX MEDLINE=95319331; PubMed=7596291;
RA Molnar I., Choi K., Yamashita M., Murooka Y.;
RT "Molecular cloning, expression in Streptomyces lividans, and analysis
of a gene cluster from Arthrobacter simplex encoding 3-
ketosteroid-DELTA.1-dehydrogenase, 3-ketosteroid-DELTA.5-isomerase
and a hypothetical regulatory protein.";
RL Mol. Microbiol. 15:895-905(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=IFO12069;
RX Dzialek J., Yamashita M., Murooka Y.;
RT "Cloning, sequencing and characterization of the downstream region of
the kddI operon of Arthrobacter simplex.";
RL Submitted (MAR-1997) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE TETR/ACRR FAMILY OF TRANSCRIPTIONAL
REGULATORS.
DR EMBL; Z93338; CAB07542.1; -.
DR InterPro; IPR001647; HTH_Tetr.
DR Pfam; PF00440; tetr; 1.
DR PRINTS; PR00455; HTH_TETR.
DR PROSITE; PS01081; HTH_TETR_FAMILY; 1.
KW DNA-binding; Hypothetical protein; Transcription regulation.
SQ SEQUENCE 212 AA; 22740 MW; F9118E18DDF4E0B2 CRC64;

Query Match 62.2%; Score 56; DB 2; Length 212;
Best Local Similarity 73.3%; Pred. No. 2.8;

Matches 11; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRRPPYLPFRPP 15
DB 83 RRRPPGSGRPP 97

RESULT 10

Q41980
ID O41980 PRELIMINARY; PRT; 212 AA.
AC O41980;
DT 01-JAN-1998 (T-Emdrel. 05, Created)
DT 01-JAN-1998 (T-Emdrel. 05, Last sequence update)
DT 01-JUN-2000 (T-Emdrel. 14, Last annotation update)
DE Hypothetical 21.9 kDa protein.
GN GAMMAHV.M13.
OS murid herpesvirus 4.
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae.
OX NCBI_TaxID=33708;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=WUMS;
RX MEDLINE=97366649; PubMed=9223479;
RA Virgin H.W. IV, Latreille P., Wamsley P., Hallsworth K., Weck K.E.,
Dal Canto A.J., Speck S.H.;
RT "Complete sequence and genomic analysis of murine gammaherpesvirus
68.";
RL J. Virol. 71:5894-5904(1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=WUMS;
RA Latreille P., Wamsley P., Waterston R.H.;
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; U97553; AAB66426.1; -.
KW Hypothetical protein.
SQ SEQUENCE 212 AA; 21911 MW; E066860064282149 CRC64;

Query Match 61.1%; Score 55; DB 12; Length 212;
Best Local Similarity 75.0%; Pred. No. 3.8;

Matches 9; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 4 PRPPYLPFRPP 15
DB 136 PSPPPLPRFP 147

RESULT 11

Q91TM2
ID Q91TM2 PRELIMINARY; PRT; 464 AA.
AC Q91TM2;
DT 01-DEC-2001 (T-Emdrel. 19, Created)
DT 01-DEC-2001 (T-Emdrel. 19, Last sequence update)
DT 01-DEC-2001 (T-Emdrel. 19, Last annotation update)
DE T74.
OS Tupaia herpesvirus.
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Betaherpesvirinae.
OX NCBI_TaxID=10397;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
RX MEDLINE=21211637; PubMed=11312357;
RA Bahr U., Darai G.;
RT "Analysis and Characterization of the Complete Genome of Tupaia (Tree
Shrew) Herpesvirus.";
RL J. Virol. 75:4854-4870(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=2;
RA Darai G., Bahr U.;
RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF281817; AAKS7119.1; -.
SQ SEQUENCE 464 AA; 51193 MW; 4BB7313EA2C2BD16 CRC64;

Query Match 61.1%; Score 55; DB 12; Length 464;
Best Local Similarity 76.9%; Pred. No. 7.7;

Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRRPPYLPFRPP 15
DB 421 RRRPPRPRPP 433

RESULT 12

Q41848
ID Q41848 PRELIMINARY; PRT; 301 AA.
AC Q41848;
DT 01-NOV-1996 (T-Emdrel. 01, Created)
DT 01-NOV-1996 (T-Emdrel. 01, Last sequence update)
DT 01-JUN-2002 (T-Emdrel. 21, Last annotation update)
DE Prolin rich protein.
GN PRP.
OS Zea mays (Maize).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade;
OC Panicoideae; Andropogoneae; Zea.
OX NCBI_TaxID=4577;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=W64A;
RX MEDLINE=92361259; PubMed=1498600;
RA Jose-Estanyol M., Ruiz-Avila L., Puigdomenech P.;
RT "A maize embryo-specific gene encodes a proline-rich and hydrophobic
protein.";
RL Plant Cell 4:413-423(1992).
DR EMBL; X60432; CAA42959.1; -.
DR HSP; P24337; 1HVP.
DR InterPro; IPR003612; AAI.
DR InterPro; IPR001768; Try/amyL inhbr.
DR Pfam; PF00234; tryp_alpha_aml; 1.
DR SMART; SM00499; AAI; 1.
SQ SEQUENCE 301 AA; 31647 MW; 884EB70854D28C2E CRC64;

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Query Match          60.6%; Score 54.5; DB 10; Length 301;
Best Local Similarity 71.4%; Pred. No. 6;
Matches 10; Conservative 1; Mismatches 2; Indels 1; Gaps 1;

QY 3 RRPYPYL-PPRPP 15
   |||: |||
Db 149 RSPYPYPPTPRPP 162

RESULT 13
P88955 ID P88955 PRELIMINARY; PRT; 2635 AA.
AC P88955;
DT 01-MAY-1997 (TREMELrel. 03, Created)
DT 01-MAY-1997 (TREMELrel. 03, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE ORF 64.
OS Kaposi's sarcoma-associated herpesvirus (KSHV) (Human herpesvirus 8).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae; Rhadinovirus.
OX NCBI_TaxID=37296;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97094384; PubMed=8939871;
RA Moore P.S., Boshoff C., Weiss R.A., Chang Y.;
RT "Molecular mimicry of human cytokine and cytokine response pathway
genes by KSHV";
RL Science 274:1739-1744 (1996).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=97121480; PubMed=8962146;
RA Russo J.J., Bohenzky R.A., Chien M.-C., Chen J., Yan M., Maddalena D.,
RA Parry J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RT "Nucleotide sequence of the Kaposi sarcoma-associated herpesvirus
(HHV8)";
RL Proc. Natl. Acad. Sci. U.S.A. 93:14862-14867 (1996).
RN [3]
RP SEQUENCE FROM N.A.
RA Russo J.J., Bohenzky R.A., Chien M.-C., Chen J., Yan M., Maddalena D.,
RA Parry J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; U75698; AAC57149.1; -.
SQ SEQUENCE 2635 AA; 289687 MW; 00070132EA8139AF CRC64;

Query Match          60.6%; Score 54.5; DB 12; Length 2635;
Best Local Similarity 68.4%; Pred. No. 44;
Matches 13; Conservative 0; Mismatches 1; Indels 5; Gaps 2;

QY 2 RRPYPYL-PPRPP 15
   |||: |||
Db 271 RRPYPYPPTPRPP 289

RESULT 14
O40942 ID O40942 PRELIMINARY; PRT; 2635 AA.
AC O40942;
DT 01-JAN-1998 (TREMELrel. 05, Created)
DT 01-JAN-1998 (TREMELrel. 05, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE ORF 64.
OS Kaposi's sarcoma-associated herpesvirus (KSHV) (Human herpesvirus 8).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae; Rhadinovirus.
OX NCBI_TaxID=37296;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97296220; PubMed=9151804;
RA Neipel F., Albrecht J.C., Fleckenstein B.;
RT "Cell-homologous genes in the Kaposi's sarcoma-associated rhadinovirus
human herpesvirus 8: determinants of its pathogenicity?";
RL J. Virol. 71:4187-4192 (1997).

[2]
RN SEQUENCE FROM N.A.
RP Neipel F., Albrecht J.-C., Ensser A., Huang Y.-Q., Li J.J.,
RA Friedman-Kien A.E., Fleckenstein B.;
RT "The genome of human herpesvirus 8 cloned from Kaposi's sarcoma.";
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; U93872; AAB62600.1; -.
SQ SEQUENCE 2635 AA; 289717 MW; 91DDA0D6FF7B660A CRC64;

Query Match          60.6%; Score 54.5; DB 12; Length 2635;
Best Local Similarity 68.4%; Pred. No. 44;
Matches 13; Conservative 0; Mismatches 1; Indels 5; Gaps 2;

QY 2 RRPYPYL-PPRPP 15
   |||: |||
Db 271 RRPYPYPPTPRPP 289

RESULT 15
Q8R353 ID Q8R353 PRELIMINARY; PRT; 347 AA.
AC Q8R353;
DT 01-JUN-2002 (TREMELrel. 21, Created)
DT 01-JUN-2002 (TREMELrel. 21, Last sequence update)
DT 01-JUN-2002 (TREMELrel. 21, Last annotation update)
DE Similar to thyroid hormone receptor-associated protein, 150 kDa
subunit.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Strausberg R.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC026580; AAH26580.1; -.
KW Receptor.
SQ SEQUENCE 347 AA; 38293 MW; C885A3C2394F4DA6 CRC64;

Query Match          60.0%; Score 54; DB 11; Length 347;
Best Local Similarity 60.0%; Pred. No. 8;
Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRPYPYPPTPRPP 15
   |||: |||
Db 262 RRPYPYPPTPRPP 276

Search completed: May 13, 2003, 10:41:27
Job time : 32 secs
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